

IN THE UNITED STATES PATENT AND TRADEMARK OFFICE

Applicant: TAMURA et al.
Title: PHARMACEUTICAL
COMPOSITIONS
Prior Appl. No.: 08/836,784
Prior Appl. Filing Date: 5/16/1997
Examiner: Unassigned
Art Unit: Unassigned

CONTINUING PATENT APPLICATION
TRANSMITTAL LETTER

Assistant Commissioner for Patents
Box PATENT APPLICATION
Washington, D.C. 20231

Sir:

Transmitted herewith for filing under 37 C.F.R. § 1.53(b) is a:

☐ Continuation ☒ Division ☐ Continuation-In-Part (CIP)

of the above-identified copending prior application in which no patenting, abandonment, or termination of proceedings has occurred. Priority to the above-identified prior application is hereby claimed under 35 U.S.C. § 120 for this continuing application. The entire disclosure of the above-identified prior application is considered as being part of the disclosure of the accompanying continuing application and is hereby incorporated by reference therein.

Enclosed are:

- ☒ Specification, Claim(s), and Abstract (61 pages).
- ☒ Declaration and Power of Attorney (2 pages).
- ☒ Preliminary Amendment.
- ☒ Information Disclosure Statement.
- ☒ Form PTO-1449.

The filing fee is calculated below:

	Claims as Filed	Included in Basic Fee	Extra Claims	Rate	Fee Totals
Basic Fee				\$690.00	\$690.00
Total Claims:	16	- 20	= 0	x \$18.00	= \$0.00
Independents:	1	- 3	= 0	x \$78.00	= \$0.00
If any Multiple Dependent Claim(s) present:			+	\$260.00	= \$0.00
				SUBTOTAL:	= \$690.00
[]				Small Entity Fees Apply (subtract ½ of above):	= \$0.00
				TOTAL FILING FEE:	= \$690.00

- [X] A check in the amount of \$690.00 to cover the filing fee is enclosed.
- [] The required filing fees are not enclosed but will be submitted in response to the Notice to File Missing Parts of Application.
- [X] The Assistant Commissioner is hereby authorized to charge any additional fees which may be required regarding this application under 37 C.F.R. §§ 1.16-1.17, or credit any overpayment, to Deposit Account No. 19-0741. Should no proper payment be enclosed herewith, as by a check being in the wrong amount, unsigned, post-dated, otherwise improper or informal or even entirely missing, the Assistant Commissioner is authorized to charge the unpaid amount to Deposit Account No. 19-0741.

Please direct all correspondence to the undersigned attorney or agent at the address indicated below.

Respectfully submitted,

Date April 18, 2000

By Patricia D. Granados

FOLEY & LARDNER
Washington Harbour
3000 K Street, N.W., Suite 500
Washington, D.C. 20007-5109
Telephone: (202) 672-5477
Facsimile: (202) 672-5399

Patricia D. Granados
Attorney for Applicant
Registration No. 33,683

IN THE UNITED STATES PATENT AND TRADEMARK OFFICE

Attorney Docket No. 087147/0437

In re patent application of

Norikazu TAMURA et al.

Serial No. Unassigned

Group Art Unit: Unassigned

Filed: April 18, 2000

Examiner: Unassigned

For: PHARMACEUTICAL COMPOSITION

Preliminary Amendment

Assistant Commissioner for Patents
Washington, D.C. 20231

Sir:

Before action is taken on the captioned application by the Examiner, Applicants respectfully request entry of the following amendments:

In the Specification:

Page 1, line 3, after the title, please insert - -This application is a divisional application of Serial No. 08/836,784, filed May 16, 1997.- -

In the Claims:

Please cancel claims 1 to 19 and 21 and amend claim 20 as follows:

20. (Amended) A method for preventing or treating angiotensin II-mediated diseases in a mammal, which comprises administering to said mammal **an effective amount of** a compound having angiotensin II antagonistic activity or a salt thereof in combination with **an effective amount of** at least one species selected from the group consisting of [a compound having the activity of increasing insulin-sensitivity,] a compound having the activity of lowering postprandial hyperglycemia in diabetes mellitus, an indane derivative having the activity of inhibiting angiotensin converting enzyme, a pyridine

derivative having the activity of inhibiting HMG-Co A reductase and their salts.

Please add the following new claims 22 to 38:

-- 22. (New) The method as claimed in Claim 20, wherein said diseases are circulatory diseases.

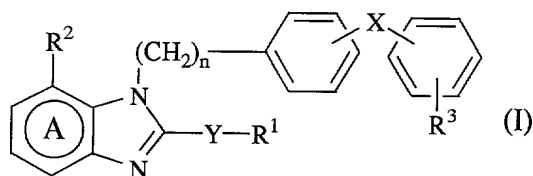
23. (New) The method as claimed in Claim 20, wherein said method is for the prevention or treatment of hypertension, cardiac insufficiency, cerebral apoplexy, ischemic peripheral circulation disturbances, myocardial ischemia, venous insufficiency, progressive cardiac insufficiency after myocardial infarction, diabetic nephropathy, nephritis, glomerulonephritis, arteriosclerosis, angiohypertrophy, vascular hypertrophy or obstruction after percutaneous transluminal coronary angioplasty, vascular reobstruction after bypass surgery, hyperaldosteronism, glomerulosclerosis, renal insufficiency, glaucoma, ocular hypertension, hyperlipemia, myocardial infarction, angina pectoris, aneurysm, coronary arteriosclerosis, cerebral arteriosclerosis, peripheral arteriosclerosis, thrombosis, diseases of central nervous system, Alzheimer's disease, deficiency of memory, depression, amnesia, senile dementia, sensory disturbances, multiple system organ failure or scleroderma, or to the prevention or amelioration of anxiety neurosis, catatonia, indisposition or dyspeptic symptoms.

24. (New) The method as claimed in Claim 20, wherein said method is for the prevention or treatment of complications of hypertension.

25. (New) The method as claimed in Claim 20, wherein said method is for the prevention or treatment of arteriosclerosis.

26. (New) The method as claimed in Claim 24, wherein said method is for the prevention or treatment of arteriosclerosis.

27. (New) The method as claimed in Claim 20, wherein the compound having angiotensin II antagonistic activity is a compound of the formula:



wherein R^1 stands for H or an optionally substituted hydrocarbon residue; R^2 stands for an optionally esterified carboxyl group; R^3 stands for a group capable of forming anion or a group convertible thereto; X shows that phenylene group and phenyl group are bonded directly or through a spacer having a chain length of 1 to 2 atoms; n denotes 1 or 2; the ring A is a benzene ring optionally having further substituents other than the group shown by R^2 ; and Y stands for a bond, -O-, -S(O)m- (m denotes 0, 1 or 2) or -N(R^4) - (R^4 stands for H or an optionally substituted alkyl group).

28. (New) The method as claimed in Claim 20, wherein the compound having angiotensin II antagonistic activity is (\pm)-1-(cyclohexyloxycarbonyloxy)ethyl 2-ethoxy-1-[[2'-(1H-tetrazol-5-yl)biphenyl-4-yl]methyl]-1H-benzimidazole-7-carboxylate, 2-ethoxy-1-[[2'-(1H-tetrazol-5-yl) biphenyl-4-yl] methyl] -1H-benzimidazole-7-carboxylic acid or 2-ethoxy-1-[[2'-(2,5-dihydro-5-oxo-1,2, 4-oxadiazol-3-yl) biphenyl-4-yl] methyl] -1H-benzimidazole-7-carboxylic acid.

29. (New) The method as claimed in Claim 20, wherein the compound having the activity of improving post-prandial hyperglycemia in diabetes mellitus is N-(1,3-dihydroxy-2-propyl) valiolamine.

30. (New) The method as claimed in Claim 20, wherein the indane derivative having the activity of inhibiting angiotensin converting enzyme is N-[N-[(S)-1-ethoxy-carbonyl-3-phenylpropyl]-L-alanyl]-N-(indan-2-yl)glycine.

31. (New) The method as claimed in Claim 20, wherein the pyridine derivative having the activity of inhibiting HMG-Co A reductase is (+)-3R,5S-erythro-(E)-7-[4-(4-fluorophenyl)-2,6-diisopropyl-5-methoxymethylpyridin-3-yl]-3,5-dihydroxyhept-6-enoic acid.

32. (New) The method as claimed in Claim 20, wherein:

(A) the compound having angiotensin II antagonistic activity is (\pm)-1-(cyclohexyloxycarbonyloxy)ethyl 2-ethoxy-1-[[2'-(1H-tetrazol-5-yl)biphenyl-4-yl]methyl]-1H-benzimidazole-7-carboxylate, 2-ethoxy-1-[[2'-(1H-tetrazol-5-yl)biphenyl-4-yl]methyl]-1H-benzimidazole-7-carboxylic acid or 2-ethoxy-1-[[2'-(2,5-dihydro-5-oxo-1,2,4-oxadiazol-3-yl)biphenyl-4-yl]methyl]-1H-benzimidazole-7-carboxylic acid;

(B) the compound having the activity of improving post-prandial hyperglycemia in diabetes mellitus is N-(1,3-dihydroxy-2-propyl) valioline;

(C) the indane derivative having the activity of inhibiting angiotensin converting enzyme is N-[N-[(S)-1-ethoxycarbonyl-3-phenylpropyl]-L-alanyl]-N-(indan-2-yl)glycine; and

(D) the pyridine derivative having the activity of inhibiting HMG-Co A reductase is (+)-3R,5S-erythro-(E)-7-[4-(4-fluorophenyl)-2,6-diisopropyl-5-methoxymethylpyridin-3-yl]-3,5-dihydroxyhept-6-enoic acid.

33. (New) The method as claimed in Claim 20, wherein the compound having angiotensin II antagonistic activity or a salt thereof is in combination with the compound having the activity of lowering postprandial hyperglycemia in diabetes mellitus or a salt thereof.

34. (New) A method as claimed in Claim 20, wherein said method is for the prevention or treatment of hypertension, arteriosclerosis or hyperlipemia and which said method comprises administering an effective amount of (\pm)-1-(cyclohexyloxycarbonyloxy)ethyl 2-ethoxy-1-[[2'-(1H-tetrazol-5-yl)biphenyl-4-yl]methyl]-1H-benzimidazole-7-carboxylate or a salt thereof in combination with an effective amount of at least one species selected from the group consisting of N-(1,3-dihydroxy-2-propyl)valioline, N-[N-[(S)-1-ethoxycarbonyl-3-phenylpropyl]-L-alanyl]-N- (indan-2-yl)glycine, (+)-3R,5S-erythro-(E)-7-[4-(4-fluorophenyl)-2,6-diisopropyl-5-methoxymethylpyridin-3-yl]-3,5-dihydroxyhept-6-enoic acid and their salts.

35. (New) A method as claimed in Claim 20, wherein said method is for the prevention or treatment of hypertension, arteriosclerosis or hyperlipemia and which said method comprises administering an effective amount of 2-ethoxy-1-[[2'-(1H-tetrazol-5-yl)biphenyl-4-yl]methyl]-1H-benzimidazole-7-carboxylic acid or a salt thereof in combination with an effective amount of at least one species selected from the group consisting of N-(1,3-dihydroxy-2-propyl)valiolamine, N-[N-[(S)-1-ethoxycarbonyl-3-phenylpropyl]-L-alanyl]-N-(indan-2-yl)glycine, (+)-3R, 5S-erythro-(E)-7-[4-(4-fluorophenyl)-2, 6-diisopropyl-5-methoxymethylpyridin-3-yl]-3,5-dihydroxyhept-6-enoic acid and their salts.

36. (New) A method as claimed in Claim 20, wherein said method is for the prevention or treatment of hypertension, arteriosclerosis or hyperlipemia and which said method comprises administering an effective amount of 2-ethoxy-1-[[2'-(2,5-dihydro-5-oxo-1,2,4-oxadiazol-3-yl)biphenyl-4-yl]methyl]-1H-benzimidazole-7-carboxylic acid or a salt thereof in combination with an effective amount of at least one species selected from the group consisting of N-(1,3-dihydroxy-2-propyl)valiolamine, N-[N-[(S)-1-ethoxycarbonyl-3-phenylpropyl]-L-alanyl]-N-(indan-2-yl)glycine, (+)-3R,5S-erythro—(E)-7-[4-(4-fluorophenyl)-2,6-diisopropyl-5-methoxymethylpyridin-3-yl]-3,5-dihydroxyhept-6-enoic acid and their salts. --

Remarks

Claims 1 to 19 and 21 have been canceled, claim 20 has been amended and claims 22 to 36 have been added. Consequently, claims 20 and 22 to 36 are pending in the captioned application. An Action on the Merits is now awaited. Should there be any questions, the Examiner is courteously invited to contact the undersigned at the telephone number shown below.

Respectfully submitted,

April 18, 2000
Date

Patricia D. Granados
Patricia D. Granados
Reg. No. 33,683

FOLEY & LARDNER
3000 K Street, N.W.
Suite 500
Washington, D.C. 20007-5109
Tel: (202) 672-5300
Fax: (202)672-5399

Should additional fees be necessary in connection with the filing of this paper, or if a petition for extension of time is required for timely acceptance of same, the Commissioner is hereby authorized to charge Deposit Account No. 19-0741 for any such fees; and applicant(s) hereby petition for any needed extension of time.

DESCRIPTION
PHARMACEUTICAL COMPOSITION

TECHNICAL FIELD

5 This invention relates to a pharmaceutical
composition comprising a compound having angiotensin II
antagonistic activity or a salt thereof in combination
with at least one species selected from the group
consisting of a compound having the activity of
10 increasing insulin-sensitivity, a compound having the
activity of improving postprandial hyperglycemia in
diabetes mellitus, an indane derivative having the
activity of inhibiting angiotensin converting enzyme, a
pyridine derivative having the activity of inhibiting
15 HMG-Co A reductase or salts of them, and to the use of
the composition.

BACKGROUND ART

Angiotensin II has a strong vasoconstrictive
action, aldosterone-synthesizing action and cell-
20 propagating action, which has been considered as one of
the mediators of various circulatory diseases. An
angiotensin II antagonistic drug suppressing the action
of angiotensin, which antagonizes to this angiotensin
II at angiotensin II receptor, is useful for the
25 prophylaxis and therapy of circulatory diseases
including hypertension, cardiac diseases (e.g. heart
failure, myocardial infarction, etc.), cerebral
apoplexy, nephritis, arteriosclerosis, etc. And, an
angiotensin converting enzyme drug suppresses
30 conversion of angiotensin I to angiotensin II, which is
considered, like angiotensin II antagonistic drugs, as
useful for the prophylaxis and therapy of circulatory
diseases including hypertension, cardiac diseases (e.g.
heart failure, myocardial infarction, etc.), cerebral
35 apoplexy, nephritis, arteriosclerosis, etc. However,
since angiotensin converting enzyme is the same enzyme

as kininase II which destructs kinin, and it has no substrate specificity, it has such an undesirable side effect as depositing inflammatory peptide including kinin and the substance P to cause occurrence of cough.

5 On the other hand, in the therapy of diabetes mellitus, there has been given treatment with a medicine to improve postprandial hyperglycemia in diabetes mellitus or treatment with a medicine to increase insulin sensitivity for preventing lowering of
10 insulin sensitivity to the intake of glucose in peripheral tissue.

Further, in the therapy of hyperlipemia, a medicine of inhibiting HMG-Co A reductase (3-hydroxy-3-methylglutaryl coenzyme A reductase) is employed to
15 suppress the biosynthesis of cholesterol.

Above all, such diseases as hypertension, abnormal carbohydrate tolerance and abnormal lipid metabolism have been known to be complicated with one another. Especially, hypertension and insulin resistance, or
20 hypertension and arteriosclerosis are considered to aggravate the respective counterpart diseases.

This invention is intended, by combination of a compound having angiotensin II antagonistic action or a salt thereof with a compound having action mechanism
25 other than the above, to perform especially remarkable effects in angiotensin II-mediated diseases, especially hypertension, hyperlipemia, arteriosclerosis and so on, singly or complications of these diseases and to cover up various defects observed in administration of a
30 medicine consisting of a single component.

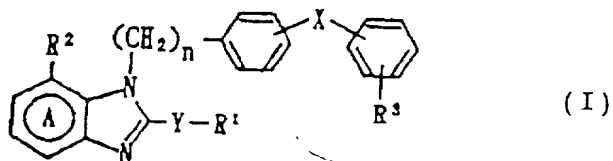
Circumstances being such as above, the present inventors have actually combined, for the first time, a compound having angiotensin antagonistic activity or a salt thereof, which is the essential component, with at
35 least one species selected from the group consisting of a compound having an insulin sensitivity increasing

action, a compound having the activity of improving
postprandial hyperglycemia in diabetes mellitus, an
indane derivative having the action of inhibiting
angiotensin converting enzyme, a pyridine derivative
5 having the action of HMG-Co A reductase or salts
thereof, and, as a result, they have found that the co-
use performs especially remarkable effects (e.g. in the
treatment effect, safety, stability, dose,
administration route, method of use, etc.) which were
10 not observed in the administration of the respective
compounds singly, and they have conducted further
studies to accomplish the present invention.

SUMMARY OF THE INVENTION

15 More specifically, the present invention relates
to
(1) a pharmaceutical composition comprising a compound
having angiotensin II antagonistic activity or a salt
thereof in combination with at least one species
selected from the group consisting of a compound having
20 the activity of increasing insulin-sensitivity, a
compound having the activity of lowering postprandial
hyperglycemia in diabetes mellitus, an indane
derivative having the activity of inhibiting
angiotensin converting enzyme, a pyridine derivative
25 having the activity of inhibiting HMG-Co A reductase
and their salts;
(2) the composition as described in the above (1),
which is a prophylactic (preventing) or therapeutic
(treating) agent of angiotensin II-mediated diseases;
30 (3) the composition as described in the above (2),
which is directed to the prophylaxis or therapy of
circulatory diseases;
(4) the composition as described in the above (2),
which is directed to the prophylaxis (prevention) or
35 therapy (treatment) of hypertension, cardiac
insufficiency, cerebral apoplexy, ischemic peripheral

- circulation disturbances, myocardial ischemia, venous insufficiency, progressive cardiac insufficiency after myocardial infarction, diabetic nephropathy, nephritis, glomerulonephritis, arteriosclerosis, angiohypertrophy, vascular hypertrophy or obstruction after percutaneous transluminal coronary angioplasty, vascular reobstruction after bypass surgery, hyperaldosteronism, glomerulosclerosis, renal insufficiency, glaucoma, ocular hypertension, hyperlipemia, myocardial infarction, angina pectoris, aneurysm, coronary arteriosclerosis, cerebral arteriosclerosis, peripheral arteriosclerosis, thrombosis, diseases of central nervous system, Alzheimer's disease, deficiency of memory, depression, amnesia, senile dementia, sensory disturbances, multiple system organ failure or scleroderma, or to the prevention or amelioration of anxiety neurosis, catatonia, indisposition or dyspeptic symptoms;
- (5) the composition as described in the above (2), which is directed to the prophylaxis or therapy of complications of hypertension;
- (6) the composition as described in the above (2), which is directed to the prophylaxis or therapy of arteriosclerosis;
- (7) the composition as described in the above (5), which is directed to the prophylaxis or therapy of arteriosclerosis;
- (8) the composition as described in the above (1), wherein the compound having angiotensin II antagonistic activity is a compound of the formula:



wherein R^1 stands for H or an optionally substituted

- hydrocarbon residue; R^2 stands for an optionally esterified carboxyl group; R^3 stands for a group capable of forming anion or a group convertible thereto; X shows that phenylene group and phenyl group are bonded directly or through a spacer having a chain length of 1 to 2 atoms; n denotes 1 or 2; the ring A is a benzene ring optionally having further substituents other than the group shown by R^2 ; and Y stands for a bond, -O-, -S(O)_m- (m denotes 0, 1 or 2) or -N(R^4)- (R^4 stands for H or an optionally substituted alkyl group);
- (9) the composition as described in the above (1), wherein the compound having angiotensin II antagonistic activity is (±)-1-(cyclohexyloxycarbonyloxy)ethyl 2-ethoxy-1-[[2'-(1H-tetrazol-5-yl)biphenyl-4-yl]methyl]-1H-benzimidazole-7-carboxylate, 2-ethoxy-1-[[2'-(1H-tetrazol-5-yl)biphenyl-4-yl]methyl]-1H-benzimidazole-7-carboxylic acid or 2-ethoxy-1-[[2'-(2,5-dihydro-5-oxo-1,2,4-oxadiazol-3-yl)biphenyl-4-yl]methyl]-1H-benzimidazole-7-carboxylic acid;
- (10) the composition as described in the above (1), wherein the compound having the activity of increasing insulin-sensitivity is 5-[4-[2-(5-ethyl-2-pyridyl)ethoxy]-benzyl]-2,4-thiazolidinedione or (R)-(+)-5-[3-[4-[2-(2-furyl)-5-methyl-4-oxazolylmethoxy]-3-methoxyphenyl]-propyl]-2,4-oxazolidinedione;
- (11) the composition as described in the above (1), wherein the compound having the activity of improving post-prandial hyperglycemia in diabetes mellitus is N-(1,3-dihydroxy-2-propyl)valiolamine;
- (12) the composition as described in the above (1), wherein the indane derivative having the activity of inhibiting angiotensin converting enzyme is N-[N-[(S)-1-ethoxy-carbonyl-3-phenylpropyl]-L-alanyl]-N-(indan-2-yl)-glycine;
- (13) the composition as described in the above (1), wherein the pyridine derivative having the activity of

inhibiting HMG-Co A reductase is (+)-3R,5S-erythro-(E)-7-[4-(4-fluorophenyl)-2,6-diisopropyl-5-methoxymethylpyridin-3-yl]-3,5-dihydroxyhept-6-enoic acid

5 [(3R,5S,6E)-7-[4-(p-fluorophenyl)-2,6-diisopropyl-5-(methoxymethyl)-3-pyridyl]-3,5-dihydroxy-6-heptenoic acid];

(14) the composition as described in the above (1), wherein the compound having angiotensin II antagonistic activity is (±)-1-(cyclohexyloxycarbonyloxy)ethyl 2-ethoxy-1-[[2'-(1H-tetrazol-5-yl)biphenyl-4-yl]methyl]-1H-benzimidazole-7-carboxylate, 2-ethoxy-1-[[2'-(1H-tetrazol-5-yl)biphenyl-4-yl]methyl]-1H-benzimidazole-7-carboxylic acid or 2-ethoxy-1-[[2'-(2,5-dihydro-5-oxo-1,2,4-oxadiazol-3-yl)biphenyl-4-yl]methyl]-1H-benzimidazole-7-carboxylic acid;

15 the compound having the activity of increasing insulin-sensitivity is 5-[4-[2-(5-ethyl-2-pyridyl)ethoxy]-benzyl]-2,4-thiazolidinedione or (R)-(+)-5-[3-[4-[2-(2-furyl)-5-methyl-4-oxazolylmethoxy]-3-methoxyphenyl]-propyl]-2,4-oxazolidinedione;

20 the compound having the activity of improving post-prandial hyperglycemia in diabetes mellitus is N-(1,3-dihydroxy-2-propyl)valiolamine;

the indane derivative having the activity of inhibiting angiotensin converting enzyme is N-[N-[(S)-1-ethoxycarbonyl-3-phenylpropyl]-L-alanyl]-N-(indan-2-yl)-glycine; and

25 the pyridine derivative having the activity of inhibiting HMG-Co A reductase is (+)-3R,5S-erythro-(E)-7-[4-(4-fluorophenyl)-2,6-diisopropyl-5-methoxymethylpyridin-3-yl]-3,5-dihydroxyhept-6-enoic acid;

30 (15) the composition as described in the above (1) comprising the compound having angiotensin II antagonistic activity or a salt thereof in combination with the compound having the activity of increasing insulin-sensitivity or a salt thereof;

(16) the composition as described in the above (1) comprising the compound having angiotensin II antagonistic activity or a salt thereof in combination with the compound having the activity of lowering postprandial hyperglycemia in diabetes mellitus or a salt thereof;

(17) a pharmaceutical composition for the prevention or treatment of hypertension, arteriosclerosis or hyperlipemia comprising (±)-1-

(cyclohexyloxycarbonyloxy)ethyl 2-ethoxy-1-[[2'-(1H-tetrazol-5-yl)biphenyl-4-yl]methyl]-1H-benzimidazole-7-carboxylate or a salt thereof in combination with at least one species selected from the group consisting of 5-[4-[2-(5-ethyl-2-pyridyl)ethoxy]benzyl]-2,4-thiazolidinedione, (R)-(+)-5-[3-[4-[2-(2-furyl)-5-methyl-4-oxazolylmethoxy]-3-methoxyphenyl]propyl]-2,4-oxazolidinedione, N-(1,3-dihydroxy-2-propyl)valiolamine, N-[N-[(S)-1-ethoxycarbonyl-3-phenylpropyl]-L-alanyl]-N-(indan-2-yl)glycine, (+)-3R,5S-erythro-(E)-7-[4-(4-fluorophenyl)-2,6-diisopropyl-5-methoxymethylpyridin-3-yl]-3,5-dihydroxyhept-6-enoic acid and their salts;

(18) a pharmaceutical composition for the prevention or treatment of hypertension, arteriosclerosis or hyperlipemia comprising 2-ethoxy-1-[[2'-(1H-tetrazol-5-yl)biphenyl-4-yl]methyl]-1H-benzimidazole-7-carboxylic acid or a salt thereof in combination with at least one species selected from the group consisting of 5-[4-[2-(5-ethyl-2-pyridyl)ethoxy]-benzyl]-2,4-thiazolidinedione, (R)-(+)-5-[3-[4-[2-(2-furyl)-5-methyl-4-oxazolylmethoxy]-3-methoxyphenyl]-propyl]-2,4-oxazolidinedione, N-(1,3-dihydroxy-2-propyl)valiolamine, N-[N-[(S)-1-ethoxycarbonyl-3-phenylpropyl]-L-alanyl]-N-(indan-2-yl)glycine, (+)-3R,5S-erythro-(E)-7-[4-(4-fluorophenyl)-2,6-diisopropyl-5-methoxymethylpyridin-3-yl]-3,5-dihydroxyhept-6-enoic acid and their salts;

(19) a pharmaceutical composition for the prevention or treatment of hypertension, arteriosclerosis or hyperlipemia comprising 2-ethoxy-1-[[2'-(1H-tetrazol-5-yl)biphenyl-4-yl]methyl]-1H-benzimidazole-7-carboxylic acid or a salt thereof in combination with at least one species selected from the group consisting of 5-[4-[2-(5-ethyl-2-pyridyl)ethoxy]-benzyl]-2,4-thiazolidinedione, (R)-(+)-5-[3-[4-[2-(2-furyl)-5-methyl-4-oxazolylmethoxy]-3-methoxyphenyl]-propyl]-2,4-oxazolidinedione, N-(1,3-dihydroxy-2-propyl)valiolamine, N-[N-[(S)-1-ethoxycarbonyl-3-phenylpropyl]-L-alanyl]-N-(indan-2-yl)glycine, (+)-3R,5S-erythro-(E)-7-[4-(4-fluorophenyl)-2,6-diisopropyl-5-methoxymethylpyridin-3-yl]-3,5-dihydroxyhept-6-enoic acid and their salts;

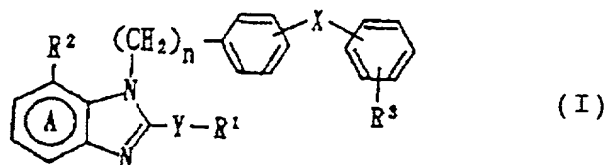
- (19) a pharmaceutical composition for the prevention or treatment of hypertension, arteriosclerosis or hyperlipemia comprising 2-ethoxy-1-[[2'-(2,5-dihydro-5-oxo-1,2,4-oxadiazol-3-yl)biphenyl-4-yl]methyl]-1H-benzimidazole-7-carboxylic acid or a salt thereof in combination with at least one species selected from the group consisting of 5-[4-[2-(5-ethyl-2-pyridyl)ethoxy]benzyl]-2,4-thiazolidinedione, (R)-(+)-5-[3-[4-[2-(2-furyl)-5-methyl-4-oxazolyl-methoxy]-3-methoxyphenyl]propyl]-2,4-oxazolidinedione, N-(1,3-dihydroxy-2-propyl)valiolamine, N-[N-[(S)-1-ethoxycarbonyl-3-phenylpropyl]-L-alanyl]-N-(indan-2-yl)glycine, (+)-3R,5S-erythro-(E)-7-[4-(4-fluorophenyl)-2,6-diisopropyl-5-methoxymethylpyridin-3-yl]-3,5-dihydroxyhept-6-enoic acid and their salts;
- (20) a method for preventing or treating angiotensin II-mediated diseases in a mammal, which comprises administering to said mammal a compound having angiotensin II antagonistic activity or a salt thereof in combination with at least one species selected from the group consisting of a compound having the activity of increasing insulin-sensitivity, a compound having the activity of lowering postprandial hyperglycemia in diabetes mellitus, an indane derivative having the activity of inhibiting angiotensin converting enzyme, a pyridine derivative having the activity of inhibiting HMG-Co A reductase and their salts; and
- (21) use of a compound having angiotensin II antagonistic activity or a salt thereof in combination with at least one species selected from the group consisting of a compound having the activity of increasing insulin-sensitivity, a compound having the activity of lowering postprandial hyperglycemia in diabetes mellitus, an indane derivative having the activity of inhibiting angiotensin converting enzyme, a pyridine derivative having the activity of inhibiting

HMG-Co A reductase and their salts, for the manufacture of a medicament for preventing or treating angiotensin II-mediated diseases.

DETAILED EXPLANATION OF THE INVENTION

Specific examples of the compound having the angiotensin II antagonistic activity or salts thereof include benzimidazol-7-carboxylic acid derivatives and salts thereof disclosed in, for example, JP-A [Japanese Patent Application Laid-open No.] H4(1992)-9373, EP-A-425921, JP-A H4(1992)-364171, EP-A-459136 and EP-A-520423, preferably compounds represented by the following formula (I) or salts thereof (preferably, pharmacologically acceptable salts).

Formula (I)



wherein R^1 stands for H or an optionally substituted hydrocarbon residue; R^2 stands for an optionally esterified carboxyl group; R^3 stands for a group capable of forming anion or a group convertible thereto; X shows that phenylene group and phenyl group are bonded directly or through a spacer having a chain length of 1 to 2 atoms; n denotes 1 or 2; the ring A is a benzene ring optionally having further substituents other than groups shown by R^2 ; and Y stands for a bond, -O-, -S(O)_m- (wherein m denotes 0, 1 or 2) or -N(R^4)- (wherein R^4 stands for H or an optionally substituted alkyl group).

In the above formula (I), examples of the hydrocarbon residue shown by R^1 include alkyl, alkenyl, alkynyl, cycloalkyl, aryl and aralkyl groups. Among them, alkyl, alkenyl and cycloalkyl groups are preferable.

The alkyl group shown by R^1 is a straight chain or branched lower alkyl group having 1 to about 8 carbon atoms, as exemplified by methyl, ethyl, propyl, isopropyl, butyl, isobutyl, sec-butyl, t-butyl, pentyl, i-pentyl, hexyl, heptyl and octyl.

The alkenyl group shown by R^1 is a straight chain or branched lower alkenyl group having 2 to about 8 carbon atoms, as exemplified by vinyl, propenyl, 2-butenyl, 3-butenyl, isobutenyl and 2-octenyl.

The alkynyl group shown by R^1 is a straight chain or branched lower alkynyl group having 2 to about 8 carbon atoms, as exemplified by ethynyl, 2-propynyl, 2-butylnyl, 2-pentylnyl and 2-octynyl.

The cycloalkyl group shown by R^1 is a lower cycloalkyl group having 3 to about 6 carbon atoms, as exemplified by cyclopropyl, cyclobutyl, cyclopentyl and cyclohexyl.

The above-mentioned alkyl, alkenyl, alkynyl or cycloalkyl group may optionally be substituted with hydroxyl group, an optionally substituted amino group (e.g. amino, N-lower (C_{1-4}) alkylamino, N,N-dilower (C_{1-4}) alkylamino, etc.), halogen, a lower (C_{1-4}) alkoxy group or a lower (C_{1-4}) alkylthio group.

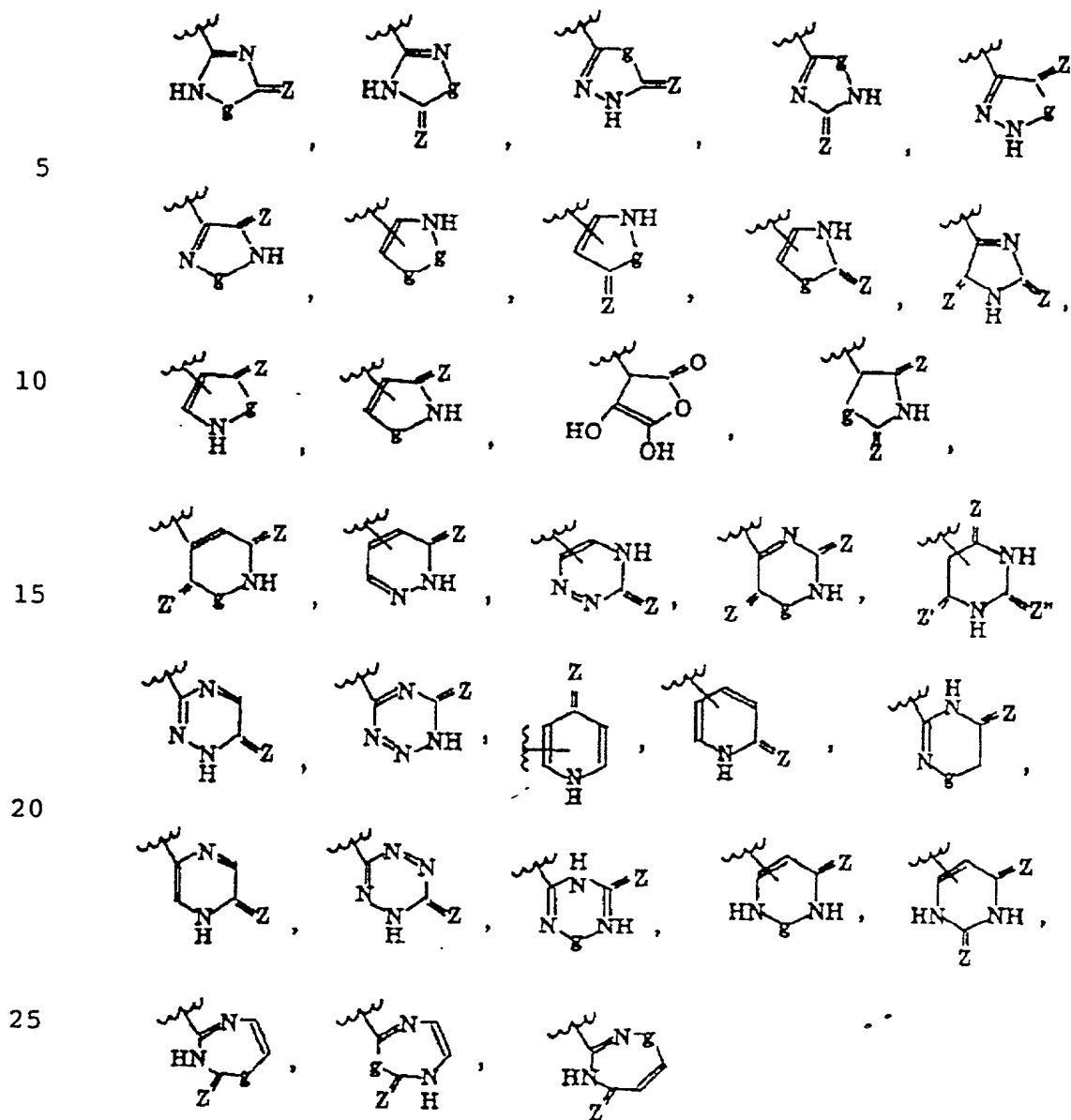
The aralkyl group shown by R^1 is exemplified by a phenyl-lower (C_{1-4}) alkyl such as benzyl, phenethyl, etc. and the aryl group shown by R^1 is exemplified by phenyl, etc.

The above-mentioned aralkyl or aryl group may optionally have, on any position of its benzene ring, for example, halogen (e.g. F, Cl, Br, etc.), nitro, an optionally substituted amino group (e.g. amino, N-lower (C_{1-4}) alkylamino, N,N-dilower (C_{1-4}) alkylamino, etc.), lower (C_{1-4}) alkoxy (e.g. methoxy, ethoxy, etc.), lower (C_{1-4}) alkylthio (e.g. methylthio, ethylthio, etc.), lower (C_{1-4}) alkyl (e.g. methyl, ethyl, etc.), etc.

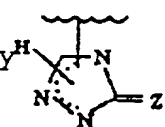
Among the above-mentioned groups shown by R^1 , optionally substituted alkyl, alkenyl or cycloalkyl groups [e.g. a lower (C_{1-5}) alkyl, lower (C_{2-5}) alkenyl or lower (C_{3-6}) cycloalkyl group optionally substituted
 5 with hydroxyl group, amino group, halogen or a lower (C_{1-4}) alkoxy group] are preferable.

Y stands for a bond, $-O-$, $-S(O)_m-$ (wherein m denotes 0, 1 or 2) or $-N(R^4)-$ (wherein R^4 stands for H or an optionally lower alkyl group), preferably a bond,
 10 $-O-$, $-S-$ or $-N(R^4)-$ [wherein R^4 stands for H or a lower (C_{1-4}) alkyl (e.g. methyl, ethyl, propyl, isopropyl, butyl, sec-butyl, t-butyl, etc.)].

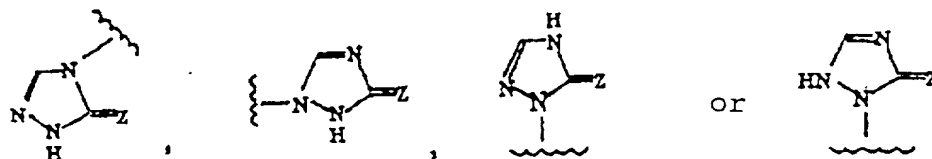
With respect to the above-mentioned formula (I), the group shown by R^3 , capable of forming anion (a
 15 group having a hydrogen atom capable of leaving as proton), or a group capable of changing thereto, is exemplified by 5- to 7-membered (preferably 5- to 6-membered) monocyclic optionally substituted heterocyclic ring residue which contain one or more of
 20 N, S and O (preferably N-containing heterocyclic ring residue having a hydrogen atom capable of leaving as proton) or groups capable of changing thereto in vivo. Such groups include the following:



30 The chemical bond between the group shown by R^3 and the partner phenyl group may be a carbon-carbon bond as shown above, or a nitrogen-carbon bond via one of the several nitrogen atoms when the symbol g stands for -NH- in the above formulae. For instance,

when R^3 is shown by , specific examples include:

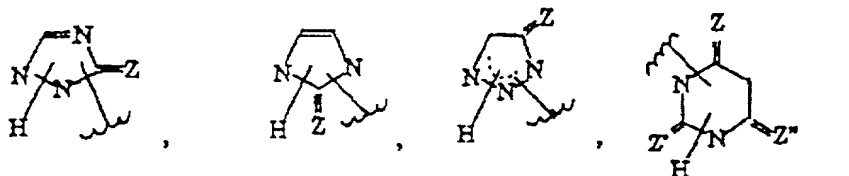
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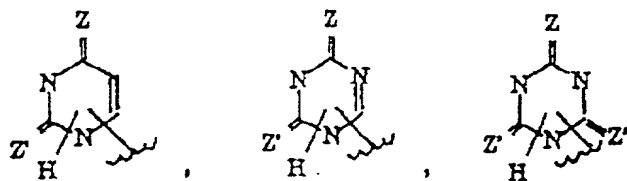
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Other examples of R^3 bonded through nitrogen atom include:

15

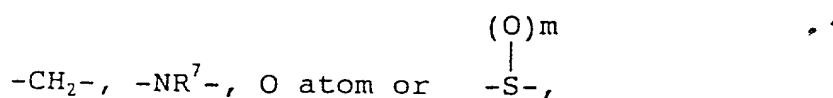


20



In the above formulae, g stands for

25



30

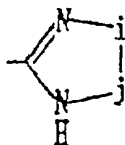
$>=\text{Z}$, $>=\text{Z}'$ and $>=\text{Z}''$ each stands for a carbonyl group, a thiocarbonyl group or an optionally oxidized sulfur atom (e.g. S, S(O), S(O)₂, etc.; preferably a carbonyl or thiocarbonyl group; more preferably, a carbonyl group); m denotes 0, 1 or 2; R^7 stands for H or an optionally substituted lower alkyl group (e.g. a lower (C₁₋₄) alkyl such as methyl, ethyl, propyl, isopropyl, butyl, sec-butyl and t-butyl).

35

Preferable examples of R^3 include 2,5-dihydro-5-

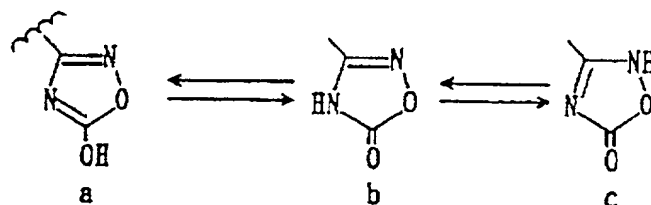
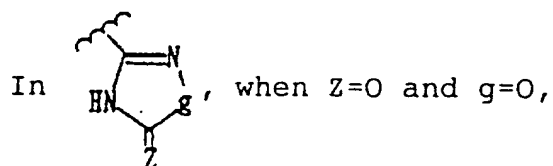
oxo-1,2,4-oxadiazole ring residue, 2,5-dihydro-5-thiooxo-1,2,4-oxadiazole ring residue or 2,5-dihydro-5-oxo-1,2,4-thiadiazole ring residue having -NH or -OH group as proton donor and carbonyl group, thiocarbonyl group or sulfinyl group as proton acceptor simultaneously.

And, while the heterocyclic residue shown by R^3 may form a condensed ring by connecting the substituents on the ring, it is preferably a 5- to 6-membered ring, more preferably a 5-membered heterocyclic residue. As R^3 , groups represented by the formula

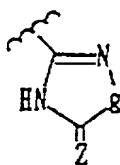


wherein i stands for -O- or -S-; j stands for $>C=O$, $>C=S$ or $>S(O)_m$; and m is of the same meaning as defined above (especially, 2,5-dihydro-5-oxo-1,2,4-oxadiazol-3-yl, 2,5-dihydro-5-thiooxo-1,2,4-oxadiazol-3-yl or 2,5-dihydro-5-oxo-1,2,4-thiadiazol-3-yl) are preferable. R^3 can be substituted at any of the ortho, meta and para position of the phenyl group, most preferably at the ortho position.

In addition, the above-mentioned heterocyclic residue (R^3) have the following tautomeric isomers. For example,



the three tautomeric isomers a, b and c exist. And, the heterocyclic residue represented by the formula



include all of the above-mentioned a, b and c.

Moreover, R^3 may be a carboxyl group, tetrazolyl group, trifluoromethanesulfonamide group ($-\text{NHSO}_2\text{CF}_3$), phosphoric acid group, sulfonic acid group, cyano group or lower (C_{1-4}) alkoxy carbonyl group; these groups each may be protected with an optionally substituted lower alkyl group or acyl group, and, any group capable of forming an anion biologically or physiologically (e.g. through biological reactions such as oxidation, reduction or hydrolysis caused by enzymes in the body) or chemically, or a group capable of changing thereto is acceptable.

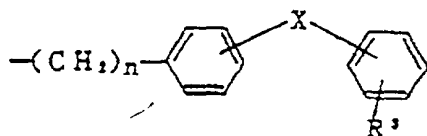
As R^3 , a tetrazolyl or carboxyl (preferably tetrazolyl) group optionally protected with an optionally substituted lower (C_{1-4}) alkyl (e.g. methyl, triphenylmethyl, methoxymethyl, ethoxymethyl, p-methoxybenzyl, p-nitrobenzyl, etc.) or acyl group (e.g. lower (C_{2-5}) alkanoyl, benzoyl, etc.) is preferable. R^3

can be substituted at any of ortho-, meta- and para-positions, preferably at the ortho-position.

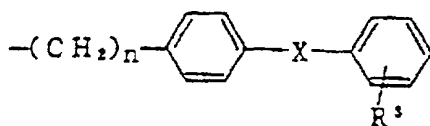
X shows the linkage of phenylene group and phenyl group adjacent to each other directly or through a spacer having a chain length of 1 to 2 atoms (preferably direct linkage). The spacer having a chain length of 1 to 2 atoms may consist of a divalent chain in which the number of atoms composing the straight chain portion is either 1 or 2, and may have a side chain, as exemplified by a lower (C_{1-4}) alkylene, $-CO-$, $-O-$, $-S-$, $-NH-$, $-CO-NH-$, $-O-CH_2-$, $-S-CH_2-$, $-CH=CH-$, etc.

The symbol n denotes an integer of 1 or 2 (preferably 1).

The formula represented by the above-mentioned R^3 , X and n:



is preferably the following one:



In the formula (I), the optionally esterified carboxyl group shown by R^2 is exemplified by groups represented by the formula $-CO-D$ [wherein D stands for a hydroxyl group or an optionally substituted alkoxy group {e.g. (i) a lower (C_{1-6}) alkoxy group whose alkyl moiety is optionally substituted with (1) a hydroxyl group, (2) an optionally substituted amino (e.g. amino, N-lower (C_{1-4}) alkylamino, N,N-lower (C_{1-4}) alkylamino, piperidino, morpholino, etc.), (3) halogen, (4) a lower (C_{1-6}) alkoxy, (5) a lower (C_{1-6}) alkylthio or (6) an optionally substituted dioxolenyl (e.g. 5-methyl-2-oxo-

1,3-dioxolen-4-yl) group, or (ii) alkoxy group shown by the formula $-O-CH(R^6)-OCOR^5$ [wherein R^6 stands for (1) H, (2) a lower (C_{1-6}) straight chain or branched alkyl group (e.g. methyl, ethyl, n-propyl, isopropyl, n-butyl, isobutyl, t-butyl, n-pentyl, isopentyl, neopentyl, etc.), (3) a lower (C_{2-6}) straight chain or branched alkenyl group (e.g. vinyl, allyl, butenyl, i-butenyl, 2-hexenyl, etc.) or (4) (C_{3-8}) cycloalkyl group (e.g. cyclopentyl, cyclohexyl, cycloheptyl, etc.); and R^5 stands for (1) a lower (C_{1-6}) straight chain or branched alkyl group (e.g. methyl, ethyl, n-propyl, isopropyl, n-butyl, isobutyl, sec-butyl, t-butyl, n-pentyl, isopentyl, neopentyl, etc.), (2) a lower (C_{2-6}) straight chain or branched alkenyl group (e.g. vinyl, allyl, butenyl, i-butenyl, 2-hexenyl, etc.), (3) a (C_{3-8}) cycloalkyl group (e.g. cyclopentyl, cyclohexyl, cycloheptyl, etc.), (4) a lower (C_{1-3}) alkyl group substituted with (C_{3-8}) cycloalkyl group (e.g. cyclopentyl, cyclohexyl, cycloheptyl, etc.) or an optionally substituted aryl group such as phenyl and naphthyl optionally substituted with halogen, nitro or a lower (C_{1-4}) alkyl (e.g. benzyl, p-chlorobenzyl, phenethyl, cyclopentylmethyl, cyclohexylmethyl, etc.), (5) a lower (C_{2-3}) alkenyl group optionally substituted with C_{3-8} cycloalkyl or an optionally substituted aryl group such as phenyl and naphthyl optionally substituted with halogen, nitro or a lower (C_{1-4}) alkyl (e.g. cinnamyl, etc. having alkenyl moiety such as vinyl, propenyl, allyl and isopropenyl), (6) an optionally substituted aryl group such as phenyl and naphthyl optionally substituted with halogen, nitro or a lower (C_{1-4}) alkyl (e.g. phenyl, p-tolyl, naphthyl, etc.), (7) a lower (C_{1-6}) straight chain or branched alkoxy group (e.g. methoxy, ethoxy, n-propoxy, isopropoxy, n-butoxy, isobutoxy, sec-butoxy, t-butoxy,

n-pentyloxy, isopentyloxy, neopentyloxy, etc.), (8) a lower (C_{2-8}) straight chain or branched alkenyloxy group (e.g. allyloxy, isobutenyloxy, etc.), (9) a (C_{3-8}) cycloalkyloxy group (e.g. cyclopentyloxy, cyclohexyloxy, cycloheptyloxy, etc.), (10) a lower (C_{1-3}) alkoxy group substituted with (C_{3-8}) cycloalkyl (e.g. cyclopentyl, cyclohexyl, cycloheptyl, etc.) or an optionally substituted aryl group such as phenyl and naphthyl optionally substituted with halogen, nitro or lower (C_{1-4}) alkyl (e.g. benzyloxy, phenethyloxy, cyclohexylmethoxy, etc. having alkoxy moiety such as methoxy, ethoxy, n-propoxy, isopropoxy, etc.), (11) a lower (C_{2-3}) lower alkenyloxy group substituted with a C_{3-8} cycloalkyl (e.g. cyclopentyl, cyclohexyl, cycloheptyl, etc.) or with an optionally substituted aryl group such as phenyl and naphthyl optionally substituted with halogen, nitro or lower (C_{1-4}) alkyl (e.g. cinnamyloxy, etc. having alkenyloxy moiety such as vinyloxy, propenyloxy, allyloxy, isopropenyloxy, etc.) or (12) an optionally substituted aryloxy group such as phenoxy and naphthoxy optionally substituted with halogen; nitro or lower (C_{1-4}) alkyl (e.g. phenoxy, p-nitrophenoxy, naphthoxy, etc.)]]]. The substituent shown by R^2 may be a group actually or potentially capable of forming anion [e.g. tetrazolyl group, trifluoromethanesulfonamide group, phosphoric acid group or sulfonic acid group optionally protected with an optionally substituted alkyl (e.g. lower (C_{1-4}) alkyl, etc.) or acyl (e.g. lower (C_{2-5}) alkanoyl, optionally substituted benzoyl, etc.)].

Examples of the substituent R^2 include -COOH and its salts, -COOMe, -COOEt, -COOtBu, -COOPr, pivaloyloxymethoxycarbonyl, 1-(cyclohexyloxycarbonyloxy)ethoxycarbonyl, (5-methyl-2-oxo-1,3-dioxolen-4-yl)methoxycarbonyl,

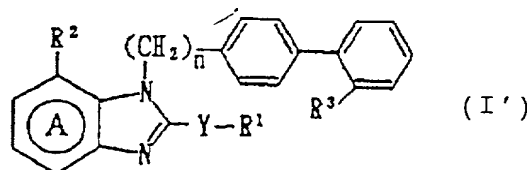
acetoxymethoxycarbonyl, propionyloxymethoxycarbonyl, n-buturyloxymethoxycarbonyl, isobutyryloxymethoxycarbonyl, (1-ethoxycarbonyloxyethoxy)carbonyl, (1-acetoxyethoxy)carbonyl, (1-isobutyryloxyethoxy)carbonyl, cyclohexylcarbonyloxymethoxycarbonyl, benzoyloxymethoxycarbonyl, cinnamyloxycarbonyl and cyclopentylcarbonyloxymethoxycarbonyl. Furthermore, R^2 may be any of the groups actually or potentially capable of forming anion (e.g. COO^- or its derivatives) under biologic or physiologic conditions (e.g. oxidation or reduction induced by enzyme present in the living body, or in vivo reaction such as hydrolysis) or chemically. R^2 may be carboxyl group or its prodrug. R^2 may be a group capable of being biologically or chemically transformed, for example, in vivo to anion.

Among the groups described above as R^2 , preferable ones include carboxyl, esterified carboxyl (e.g. methyl ester, ethyl ester or an ester formed by binding of a group shown by the formula $-\text{O}-\text{CH}(\text{R}^6)-\text{OCOR}^5$ to carbonyl) and optionally protected tetrazolyl, carboaldehyde and hydroxymethyl.

In the formula (I), ring A may have, in addition to the group shown by R^2 , further substituents as exemplified by halogen (e.g. F, Cl, Br, etc.), cyano, nitro, lower (C_{1-4}) alkyl, lower (C_{1-4}) alkoxy, an optionally substituted amino group {e.g. amino, N-lower (C_{1-4}) alkylamino (e.g. methylamino, etc.), N,N-di-lower (C_{1-4}) alkylamino (e.g. dimethylamino, etc.), N-arylamino (e.g. phenylamino, etc.), alicyclic amino (e.g. morpholino, piperidino, piperazino, N-phenylpiperazino, etc.)}, a group shown by the formula $-\text{CO}-\text{D}'$ [wherein D' stands for hydroxyl group or

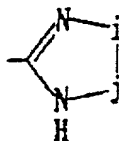
a lower (C₁₋₄) alkoxy group whose alkyl moiety may optionally be substituted with hydroxyl group, lower (C₁₋₄) alkoxy, lower (C₂₋₆) alkanoyloxy (e.g. acetoxy, pivaloyloxy, etc.) or lower (C₁₋₆) alkoxycarbonyloxy (e.g. chain-like alkoxycarbonyloxy such as methoxycarbonyloxy, ethoxycarbonyloxy, etc. or cyclic alkoxycarbonyloxy such as cyclohexyloxycarbonyloxy)], or a tetrazolyl group, a trifluoromethanesulfonamide group, a phosphoric acid group or a sulfonic acid group which may optionally be protected with lower (C₁₋₄) alkyl or acyl (e.g. lower (C₂₋₅) alkanoyl, optionally substituted benzoyl, etc.); among them, a lower (C₁₋₄) alkyl and halogen are preferable. Of these substituents, one or two may simultaneously be substituted at available positions in the ring.

Among the compounds represented by the formula (I) mentioned above, compounds represented by the formula (I') or salts thereof are preferred:



wherein ring A stands for a benzene ring optionally having further substituents besides groups shown by R²; R¹ stands for H or an optionally substituted lower (C₁₋₆) alkyl (preferably lower alkyl (C₁₋₄) alkyl); Y stands for -O-, -S- or -N(H)-; R² is a group represented by the formula -CO-D" [wherein D" stands for hydroxyl group, or a lower (C₁₋₄) alkoxy whose alkyl moiety is optionally substituted with hydroxyl group, amino, halogen, a lower (C₂₋₆) alkanoyloxy (e.g. acetyloxy, pivaloyloxy, etc.), a lower (C₄₋₇) cycloalkanoyloxy, (lower (C₁₋₆) alkoxy) carbonyloxy (e.g. methoxycarbonyloxy, ethoxycarbonyloxy, etc.), (lower

(C₃₋₇)cycloalkoxy)carbonyloxy (e.g. cyclohexyloxycarbonyl, etc.) or a lower (C₁₋₄)alkoxy; R³ stands for a tetrazolyl, carboxyl group or groups represented by the formula



[wherein i stands for -O- or -S-; j stands for >C=O, >C=S or >S(O)_m, m denotes 0, 1 or 2] each of which is optionally protected with optionally substituted lower (C₁₋₄) alkyl (e.g. methyl, triphenylmethyl, methoxymethyl, acetyloxymethyl, methoxycarbonyloxymethyl, ethoxycarbonyloxymethyl, 1-(cyclohexyloxycarbonyloxy)ethyl, pivaloyloxymethyl, etc.) or an acyl group (e.g. a lower (C₂₋₅) alkanoyl, benzoyl, etc.); n denotes 1 or 2 (preferably 1)].

In the formula (I'), as substituents on the optionally substituted lower alkyl shown by R¹, mention is made of a hydroxyl group, an amino group, halogen or a lower (C₁₋₄) alkoxy group.

In the formula (I'), as substituents other than those shown by R² on the ring A, mention is made of halogen (e.g. F, Cl, Br, etc.), lower (C₁₋₄) alkyl, lower (C₁₋₄) alkoxy, nitro, a group represented by the formula -CO-D' [wherein D' stands for a hydroxyl group or lower (C₁₋₄) alkoxy whose alkyl moiety may optionally be substituted with hydroxyl group, lower (C₁₋₄) alkoxy, lower (C₂₋₆) alkanoyloxy (e.g. acetoxy, pivaloyloxy, etc.) or lower (C₁₋₆) alkoxycarbonyloxy (e.g. methoxycarbonyloxy, ethoxycarbonyloxy, cyclohexyloxycarbonyloxy, etc.)] or amino optionally substituted with a lower (C₁₋₄) alkyl (preferably lower (C₁₋₄) alkyl or halogen). As the ring A, a benzene ring, which has no substituent other than the group

represented by the formula R^2 , is more preferable.

As the salts mentioned above, mention is made of pharmaceutically acceptable ones, as exemplified by a salt with an inorganic base, an organic base, an inorganic acid, an organic acid, or a basic or acidic amino acid. Preferable examples of a salt with an inorganic base include alkali metal salts such as sodium salts, potassium salts, and so on; alkaline earth metal salts such as calcium salts, magnesium salts, and so on; as well as aluminum salts, ammonium salts, etc. Preferable examples of a salt with an organic base include salts with trimethylamine, triethylamine, pyridine, picoline, ethanolamine, diethanolamine, triethanolamine, dicyclohexylamine, N,N'-dibenzylethylenediamine, N-methylmorpholine, etc. Preferable examples of a salt with an inorganic acid include salts with hydrochloric acid, hydrobromic acid, nitric acid, sulfuric acid, phosphoric acid, etc. Preferable examples of the salt with an organic acid include salts with formic acid, acetic acid, trifluoroacetic acid, fumaric acid, oxalic acid, tartaric acid, maleic acid, citric acid, succinic acid, malic acid, methanesulfonic acid, benzenesulfonic acid, p-toluenesulfonic acid, etc. Preferable examples of a salt with a basic amino acid include salts with arginine, lysine, ornithine, etc. Preferable examples of a salt with an acidic amino acid include salts with aspartic acid, glutamic acid, etc.

Preferable compounds to be employed as the active ingredient of the present invention include those described in the Examples of JP-A H4(1992)-364171/1992, EP-A-459136 and EP-A-520423. Among them, (\pm)-1-(cyclohexyloxycarbonyloxy)ethyl 2-ethoxy-1-[[2'-(1H-tetrazol-5-yl)biphenyl-4-yl]methyl]-1H-benzimidazole-7-carboxylate, 2-ethoxy-1-[[2'-(1H-tetrazol-5-yl)biphenyl-4-yl]methyl]-1H-benzimidazole-7-carboxylic

acid, 2-ethoxy-1-[[2'-(2,5-dihydro-5-oxo-1,2,4-oxadiazol-3-yl)biphenyl-4-yl]methyl]-1H-benzimidazole-7-carboxylic acid or pharmaceutically acceptable salts thereof are preferable.

5 The compounds represented by the general formula (I) are disclosed in, for example, JP-A H4(1992)-9373, EP-A-425921, JP-A H4(1992)-364171, EP-A-459136 and EP-A-520423, which can be produced by the methods disclosed in these official publications or methods
10 analogous thereto.

 As the compound having the activity of increasing insulin-sensitivity to be used for the present invention or salts thereof, mention is made of a compound having the activity of normalizing the
15 function of the receptor whose insulin-activity is damaged, namely a compound having the activity of releasing the insulin-resistance, or salts thereof. Specific examples of such compounds as above include 2,4-thiazolidinedione, 2,4-oxazolidinedione derivatives
20 or salts thereof described in EP-A-193256, Japan Patent Application No. H7(1995)-284106 (EP-A-710659), JP-A S60(1985)-51189, or known compounds having the activity of increasing insulin-sensitivity, for example, 5-[[3,4-dihydro-2-(phenylmethyl)-2H-1-benzopyran-6-yl]methyl]-2,4-thiazolidinedione (generic name:
25 englitazone); 5-[[4-[3-(5-methyl-2-phenyl-4-oxazolyl)-1-oxopropyl]phenyl]methyl]-2,4-thiazolidinedione (generic name: darglitazone; CP-86325);
30 5-[2-(5-methyl-2-phenyl-4-oxazolylmethyl)benzofuran-5-ylmethyl]-2,4-oxazolidinedione (CP-92768); 5-(2-naphthalenylsulfonyl)-2,4-thiazolidinedione (AY-31637);
 4-[(2-naphthalenyl)methyl]-3H-1,2,3,5-oxathiadiazol-2-oxide (AY-30711); and
35 5-[[4-[2-(methyl-2-pyridylamino)ethoxy]phenyl]methyl]-

2,4-thiazolidinedione (BRL-49653). Preferable compounds include those described as Working Examples in EP-A-193256, Japan Patent Application No. H7(1995)-284106 (EP-A-710659) or JP-A S60(1985)-51189. Among them,
 5 2,4-thiazolidinedione or 2,4-oxazolidinedione derivatives such as 5-[4-[2-(3-ethyl-2-pyridyl)ethoxy]benzyl]-2,4-thiazolidinedione, 5-[4-[2-(5-ethyl-2-pyridyl)ethoxy]benzyl]-2,4-thiazolidinedione, 5-[4-[2-(5-ethyl-2-pyridyl)ethoxy]benzyl]-2,4-thiazolidinedione, 5-[4-[2-(5-ethyl-2-pyridyl)ethoxy]benzyl]-2,4-thiazolidinedione, (R)-(+)-5-[3-[4-[2-(2-furyl)-5-methyl-4-oxazolylmethoxy]-3-methoxyphenyl]propyl]-2,4-oxazolidinedione and CS-045
 10 are preferable, especially, 5-[4-[2-(5-ethyl-2-pyridyl)ethoxy]benzyl]-2,4-thiazolidinedione or (R)-(+)-5-[3-[4-[2-(2-furyl)-5-methyl-4-oxazolylmethoxy]-3-methoxyphenyl]propyl]-2,4-oxazolidinedione is preferable.

Preferable examples of salts of a compound having
 20 the activity of increasing (enhancing) insulin-sensitivity include pharmaceutically acceptable salts, which are specifically exemplified by substantially the same ones as pharmaceutically acceptable salts of the above-mentioned compounds having the angiotensin II
 25 antagonistic activity.

As the compound having the activity of improving postprandial hyperglycemia in diabetes mellitus or salts thereof to be used in the present invention, mention is made of a compound having the activity of
 30 inhibiting α -glucosidase and having the activity of inhibiting a digestive enzyme such as amilase, maltase, α -dextrinase, sucrase and so on to delay the digestion of starch or sucrose, or salts thereof. As examples of them, mention is made of valiolumine derivatives or
 35 salts thereof described in EP-A-56194, etc., acarbose or salts thereof described in USP 4062950, etc. As

preferable examples of them, mention is made of compounds described in Examples of EP-A-56194, and, among them, N-(1,3-dihydroxy-2-propyl)valiolamine is preferable.

5 Preferable examples of salts of a compound having the activity of improving postprandial hyperglycemia in diabetes mellitus include pharmaceutically acceptable salts, which are specifically exemplified by substantially the same ones as pharmaceutically
10 acceptable salts of the above-mentioned compounds having the angiotensin II antagonistic activity.

As indane derivatives having the activity of inhibiting angiotensin converting enzyme or salts thereof to be used in the present invention, mention is
15 made of indane derivatives or salts thereof having the antihypertensive activity by inhibiting angiotensin converting enzyme which converts angiotensin I to angiotensin II. Specific examples of them include indane derivatives or salts thereof described in, for
20 example, JP-A S57(1982)-179141 and EP-A-51391. As preferable compounds, mention is made of those described as Working Examples in JP-A S57(1982)-179141 or EP-A-51391. Among them, N-[N-[(S)-1-ethoxycarbonyl-3-phenylpropyl]-L-alanyl]-N-(indan-2-yl)glycine or
25 salts thereof are preferable, and especially, N-[N-[(S)-1-ethoxycarbonyl-3-phenylpropyl]-L-alanyl]-N-(indan-2-yl)glycine hydrochloride is preferable.

As preferable examples of salts of indane derivatives having the activity of inhibiting
30 angiotensin converting enzyme, mention is made of pharmaceutically acceptable salts. As specific examples of them, mention is made of those which are substantially the same as pharmaceutically acceptable salts of the above-mentioned compound having the
35 angiotensin II antagonistic activity.

In the present invention, a compound having the

angiotensin II antagonistic activity or a salt thereof is used in combination with an indane derivative having the activity of inhibiting angiotensin converting enzyme or a salt thereof. In place of the above-mentioned indane derivative having the activity of inhibiting angiotensin converting enzyme, other angiotensin converting enzyme inhibiting agents (e.g. captopril, enalapril, alacepril, ramipril, lisinopril imidapril, etc.) may optionally be used, and, any other antihypertensive agent such as α -blocker, β -blocker, a diuretic or a calcium antagonist may optionally be used in combination with an angiotensin II antagonist.

As the pyridine derivative having the activity of inhibiting HMG-Co A reductase or a salt thereof to be used in the present invention, mention is made of a pyridine derivative having the activity of inhibiting HMG-Co A reductase, which is a rate-limiting enzyme of cholesterol synthesis, or a salt thereof. Specific example of them include pyridine derivatives or salts thereof described in, for example, JP-A H1(1989)-216974, EP-A-325130, JP-A H4(1992)-308573, USP 5177080, JP-B [Japanese Patent Examined Publication No.] H6(1994)-41448, EP-A-307342, JP-A H1(1989)-121266 and EP-A-306929. As preferable compounds, mention is made of, for example, pyridine derivatives described as Working Examples in these official publications, and, among them, pyridine derivatives described as Working Examples in JP-A H4(1992)-308573 are more preferable, especially preferable one being (+)-3R,5S-erythro-(E)-7-[4-(4-fluorophenyl)-2,6-diisopropyl-5-methoxymethylpyridin-3-yl]-3,5-dihydroxyhept-6-enoic acid or salts thereof and most preferable one being (+)-3R,5S-erythro-(E)-7-[4-(4-fluorophenyl)-2,6-diisopropyl-5-methoxymethylpyridin-3-yl]-3,5-dihydroxyhept-6-enoate sodium.

As preferable examples of salts of a pyridine

derivative having the activity of inhibiting HMG-Co A reductase, mention is made of pharmaceutically acceptable salts, which are specifically exemplified by substantially the same ones as pharmaceutically acceptable salts of the above-mentioned angiotensin II antagonistic compounds.

In the present invention, an angiotensin II antagonistic compound or a salt thereof is used in combination with a pyridine derivative having the activity of inhibiting HMG-Co A reductase or a salt thereof. And, in place of the above-mentioned pyridine derivatives having the activity of inhibiting HMG-Co A reductase, any other agent of inhibiting HMG-Co A reductase (e.g. pravastatin, simvastatin, lovastatin or fluvastatin may optionally be employed. And, any other antihyperlipemic drug including an agent of inhibiting squalene synthesis and a fibrate compound having the activity of lowering triglyceride (e.g. bezafibrate) may optionally be used in combination with an angiotensin II antagonistic drug.

In the present invention, a compound having the angiotensin II antagonistic activity or a salt thereof is employed in combination with at least one species of a compound having the activity of increasing insulin-sensitivity, a compound having the activity of improving postprandial hyperglycemia in diabetes mellitus, an indane derivative having the activity of inhibiting angiotensin converting enzyme, a pyridine derivative having the activity of inhibiting HMG-Co A reductase or salts thereof. And, a combination of one or more species of a compound having the activity of increasing insulin-sensitivity, a compound having the activity of improving postprandial hyperglycemia in diabetes mellitus, an indane derivative having the activity of inhibiting angiotensin converting enzyme, a pyridine derivative having the activity of inhibiting

HMG-Co A reductase or salts of them may optionally be employed. And, any other drugs (e.g. an antihypertensive drug, an antihyperlipemic drug, etc.) may optionally be combined appropriately with any one of the above compound.

To state further, in the case of using a compound having the angiotensin antagonistic activity or a salt thereof in combination with at least one species of a compound having the activity of increasing insulin-sensitivity, a compound having the activity of improving postprandial hyperglycemia in diabetes mellitus, an indane derivative having the activity of inhibiting angiotensin converting enzyme, a pyridine derivative having the activity of inhibiting HMG-Co A reductase or salts thereof, these drugs can be formulated by mixing individually or simultaneously with pharmaceutically acceptable carriers, excipients, binders, diluents or the like, which can be administered orally or non-orally. In the case of formulating these effective components individually, while thus individually formulated agents can be administered in the form of their mixture prepared by using, for example, a diluent when administered, the individually formulated agents can also be administered separately or simultaneously or with time intervals to the one and same subject. A kit for administering the individually formulated effective components in the form of their mixture prepared by using, for example, a diluent when administered (e.g. a kit for injection which comprises two or more ampoules each comprising a powdery component and a diluent for mixing and dissolving two or more components when administered, etc.), a kit for administering the individually formulated agents simultaneously or with time intervals to the one and same subject (e.g. a kit for tablets to be administered simultaneously or with time intervals,

characterized by having two or more tablets each comprising an agent and said tablets being put in one or separate bags and, if necessary, a column to describe time to be administered each agent, etc.) are also included by the pharmaceutical composition of the present invention.

Preferable combinations of the pharmaceutical composition of the present invention are as follows:

- (1) a combination of (\pm)-1-(cyclohexyloxycarbonyloxy)ethyl 2-ethoxy-1-[[2'-(1H-tetrazol-5-yl)biphenyl-4-yl]methyl]-1H-benzimidazole-7-carboxylate or a salt thereof with at least one species of 5-[4-[2-(5-ethyl-2-pyridyl)ethoxy]benzyl]-2,4-thiazolidinedione, (R)-(+)-5-[3-[4-[2-(2-furyl)-5-methyl-4-oxazolylmethoxy]-3-methoxyphenyl]propyl]-2,4-oxazolidinedione, N-(1,3-dihydroxy-2-propyl)valiolamine, N-[N-[(S)-1-ethoxycarbonyl-3-phenylpropyl]-L-alanyl]-N-(indan-2-yl)glycine, (+)-3R,5S-erythro-(E)-7-[4-(4-fluorophenyl)-2,6-diisopropyl-5-methoxymethylpyridin-3-yl]-3,5-dihydroxyhept-6-enoic acid or salts thereof;
- (2) a combination of 2-ethoxy-1-[[2'-(1H-tetrazol-5-yl)biphenyl-4-yl]methyl]-1H-benzimidazole-7-carboxylic acid or a salt thereof with at least one species of 5-[4-[2-(5-ethyl-2-pyridyl)ethoxy]benzyl]-2,4-thiazolidinedione, (R)-(+)-5-[3-[4-[2-(2-furyl)-5-methyl-4-oxazolylmethoxy]-3-methoxyphenyl]propyl]-2,4-oxazolidinedione, N-(1,3-dihydroxy-2-propyl)valiolamine, N-[N-[(S)-1-ethoxycarbonyl-3-phenylpropyl]-L-alanyl]-N-(indan-2-yl)glycine, (+)-3R,5S-erythro-(E)-7-[4-(4-fluorophenyl)-2,6-diisopropyl-5-methoxymethylpyridin-3-yl]-3,5-dihydroxyhept-6-enoic acid or salts thereof, and
- (3) a combination of 2-ethoxy-1-[[2'-(2,5-dihydro-5-oxo-1,2,4-oxadiazol-3-yl)biphenyl-4-yl]methyl]-1H-benzimidazole-7-carboxylic acid or a salt thereof with

at least one species of 5-[4-[2-(5-ethyl-2-pyridyl)ethoxy]benzyl]-2,4-thiazolidinedione, (R)-(+)-5-[3-[4-[2-(2-furyl)-5-methyl-4-oxazolylmethoxy]-3-methoxyphenyl]propyl]-2,4-oxazolidinedione, N-(1,3-

5 dihydroxy-2-propyl)valiolamine, N-[N-[(S)-1-ethoxycarbonyl-3-phenylpropyl]-L-alanyl]-N-(indan-2-yl)glycine, (+)-3R,5S-erythro-(E)-7-[4-(4-fluorophenyl)-2,6-diisopropyl-5-methoxymethylpyridin-

yl]-3,5-dihydroxyhept-6-enoic acid or salts thereof.

10 These preferred combinations (1) to (3) are preferably used for the prevention or treatment of hypertension, arteriosclerosis or hyperlipemia, in particular, arteriosclerosis accompanied with hypertension.

Among them, a combination of a compound having the

15 angiotensin II antagonistic activity or a salt thereof with at least one species of a compound having the activity of increasing insulin-sensitivity, a compound having the activity of improving postprandial hyperglycemia in diabetes mellitus or salts of them is

20 preferably used.

The pharmaceutical composition of this invention is used as a prophylactic or therapeutic agent of, for example, angiotensin II-mediated diseases of animals, especially mammals (e.g. man, dog, rabbit, rat, mouse,

25 etc.), as exemplified by circulatory diseases including hypertension, cardiac insufficiency, cerebral apoplexy, ischemic peripheral circulation disturbances, myocardial ischemia, venous insufficiency, progressive cardiac insufficiency after myocardial infarction,

30 diabetic nephropathy, nephritis, glomerulonephritis, arteriosclerosis, angiohypertrophy, vascular hypertrophy or obstruction after percutaneous transluminal coronary angioplasty, vascular reobstruction after bypass surgery,

35 hyperaldosteronism, glomerulosclerosis, renal insufficiency, glaucoma, ocular hypertension,

hyperlipemia, myocardial infarction, angina pectoris, aneurysm, coronary arteriosclerosis, cerebral arteriosclerosis, peripheral arteriosclerosis, thrombosis; diseases of sensory disturbances including
5 Alzheimer's disease, deficiency of memory, depression, amnesia, senile dementia; diseases of central nervous system including anxiety neurosis, catatonia and indisposition; dyspeptic symptoms, multiple system organ failure, and scleroderma. The pharmaceutical
10 composition of this invention is preferably used as a prophylactic or therapeutic agent for, especially, circulatory diseases including diseases of central nervous system caused by circulatory disturbances. Among the circulatory diseases, for the prophylaxis or
15 therapy of arteriosclerosis and hyperlipemia, use of the pharmaceutical composition of this invention is preferable, especially, use of it for the prophylaxis or therapy of arteriosclerosis is preferable. Further, also for the therapeutic method for lowering
20 cholesterol, the pharmaceutical composition of this invention can be used.

And, the pharmaceutical composition of this invention performs remarkable effects for the prophylaxis or therapy of diseases accompanied with
25 diabetic, obesitic, hyperlipemic or essential hypertension. It is preferably used, especially, for the prophylaxis or therapy of arteriosclerosis accompanied with hypertension.

The pharmaceutical composition of this invention
30 can be administered orally or non-orally in the form of, for example, granules, powdery preparations, dust preparations, tablets, capsules, syrup, emulsions, suppositories (e.g. rectal suppositories and vaginal
35 suppositories), injections (e.g. subcutaneous, intravenous, intramuscular or intraperitoneal injections), instillation, medicines for external

application (e.g. preparations to be administered through nasal route, transdermally administrable preparations and ointments), emulsions, elixir, suspensions and solutions. These preparations can be formulated in accordance with per se known methods usually employed in the formulation process. In the present specification, the term "non-orally" includes subcutaneous injection, intravenous injection intramuscular injection, intraperitoneal injection or instillation.

Injectable preparations, for example, sterile injectable aqueous suspensions or oil suspensions can be prepared by known procedures in the relevant fields, using a suitable dispersant or wetting agent and suspending agent. The sterile injections may be in the state of, for example, a solution or a suspension, which is prepared with a non-toxic diluent administrable non-orally, e.g. an aqueous solution, or with a solvent employable for sterile injection. Examples of usable vehicles or acceptable solvents include water, Ringer's solution and an isotonic aqueous saline solution. Further, a sterile non-volatile oil can usually be employed as solvent or suspending agent.

Any non-volatile oil and a fatty acid can be used for this purpose, which include natural or synthetic or semi-synthetic fatty oil or fatty acid, and natural or synthetic or semi-synthetic mono- or di- or tri-glycerides.

Furthermore, additives including a preservative, an isotonizer, a solubilizer, a stabilizer and a pain-soothing agent may adequately be employed.

Rectal suppositories can be prepared by mixing the drug with a suitable non-irritable vehicle, for example, cocoa butter and polyethylene glycols, which are in the solid state at ordinary temperatures, but,

in the liquid state at temperatures in intestinal tubes and melt in rectum to release the drug.

As a solid formulation for oral administration, mention is made of powdery preparations, granules, tablets, pills and capsules as referred to in the above. In such formulations as exemplified above, the active components can be mixed with at least one additive, for example, sucrose, lactose, cellulose sugar, mannitol, maltitol, dextran, starch, agar, alginates, chitins, chitosans, pectins, tragacanth gum, gum arabic, gelatins, collagens, casein, albumin, synthetic or semi-synthetic polymers or glycerides. These formulations can contain, as in conventional cases, further additives, for example, an inactive diluent, a lubricant such as magnesium stearate, a preservative such as parabens and sorbic acid, an antioxidant such as ascorbic acid, α -tocopherol or cysteine, an excipient, a disintegrator, a binder, a thickening agent, a buffer, a sweetener, a flavoring agent, a perfuming agent and a coating agent. Tablets and pills can further be prepared with enteric coating. Examples of liquid preparations for oral administration include pharmaceutically acceptable emulsions, syrups, elixirs, suspensions and solutions, which may contain an inactive diluent, for example, water, which is conventionally employed in the relevant field.

A formulation used for the pharmaceutical composition of this invention preferably comprises, as an effective component, about 0.6 to 39 weight % (more preferably about 0.7 to 27 weight %) of a compound having angiotensin II antagonistic activity or a salt thereof, about 0.06 to 35 weight % (more preferably about 0.6 to 23 weight %) of a compound having the activity of increasing insulin-sensitivity or a salt thereof, about 0.06 to 0.39 weight % (more preferably about 0.06

to 0.24 weight %) of a compound having the activity of improving postprandial hyperglycemia in diabetes mellitus or a salt thereof,

5 about 3 to 46 weight % (more preferably about 3 to 23 weight %) of an indane derivative having the activity of inhibiting angiotensin converting enzyme or a salt thereof and/or
about 0.006 to 0.77 weight % (more preferably about 0.006 to 0.39 weight %) of a pyridine derivative having
10 the activity of inhibiting HMG-Co A reductase or salt thereof.

This formulation may be prepared by formulating two or more components individually or simultaneously.

15 The pharmaceutical composition of this invention is less toxic, which is safely used for animals, especially mammals (e.g. man, dog, rabbit, rat, mouse, etc.) and can be used advantageously for prophylaxis or therapy of angiotensin II-mediated diseases.

20 The dose of the pharmaceutical composition of this invention is determined in accordance with the dose of individual drugs, and can be selected dependent on the age, body weight, symptom, dose interval, administration routes, type of the formulation, and combination of drugs.

25 The dose to be administered to a specific patient is dependent on the age, body weight, general health conditions, sex, diet, dose interval, administration routes, excretion rate, combination of drugs and conditions of the diseases then treated, while taking
30 the minimal recommendable clinical dose or these and other necessary factors into consideration.

Typical daily doses of the compositions having various combinations of an angiotensin II antagonistic compound or a salt thereof with at least one species of
35 a compound having the activity of increasing insulin-sensitivity, a compound having the activity of

improving postprandial hyperglycemia in diabetes mellitus, an indane derivative having the activity of inhibiting angiotensin converting enzyme, a pyridine derivative having the activity of inhibiting HMG-Co A reductase and salts thereof are within the range of from about 1/50 of the minimal recommendable clinical dose to maximal recommendable dose (preferably minimum recommendable dose, more preferably about 1/2 of minimum recommendable dose) in the case of practical administration of these compounds individually.

For example, in case of the treatment of arteriosclerosis in human adult (body weight: about 60 kg), (\pm)-1-(cyclohexyloxycarbonyloxy)ethyl 2-ethoxy-[[2'-(1H-tetrazol-5-yl)biphenyl-4-yl]methyl]-1H-benzimidazole-7-carboxylate in a dose ranging from about 1 to 50 mg/patient/day (preferably from about 1 to 35 mg/patient/day) can be effectively combined with, for example, 5-[4-[2-(5-ethyl-2-pyridyl)ethoxy]benzyl]-2,4-thiazolidindione in a dose ranging from about 0.1 to 30 mg/patient/day (preferably from about 2 to 30 mg/patient/day) or N-(1,3-dihydroxy-2-propyl)valiolamine in a dose ranging from about 0.1 to 2 mg/patient/day. Needless to state, while these dosage ranges can be adjusted by a necessary unit base for dividing a daily dose, such doses are decided by taking into consideration the diseases to be treated, conditions of such diseases, the age, body weight, general health conditions, sex and diet of the patient then treated, dose internals, administration routes, excretion rate, combinations of drugs or any other necessary factors into consideration. In the prophylactic or therapeutic agents of this invention, the unit dose is administered once or twice daily (preferably once).

In case of the prevention or treatment of arteriosclerosis of human adult (body weight: about 60

kg), preferred embodiments of the above-mentioned preferred combinations (1) to (3) are shown below:

(1) A tablet comprising about 1 to 50 mg (preferably about 1 to 35 mg) of (+)-1-

5 (cyclohexyloxycarbonyloxy)ethyl 2-ethoxy-1-[[2'-(1H-tetrazol-5-yl)biphenyl-4-yl]methyl]-1H-benzimidazole-7-carboxylate is orally administered to one and same subject in the form of combination use with a tablet comprising about 0.1 to 45 mg (preferably about 2 to 30
10 mg) of 5-[4-[2-(5-ethyl-2-pyridyl)ethoxy]benzyl]-2,4-thiazolidinedione, a tablet comprising about 1 to 20 mg (preferably about 1 to 15 mg) of (R)-(+)-5-[3-[4-[2-(2-furyl)-5-methyl-4-oxazolylmethoxy]-3-methoxyphenyl]propyl]-2,4-oxazolidinedione, a tablet
15 comprising about 0.1 to 0.5 mg (preferably about 0.1 to 0.3 mg) of N-(1,3-dihydroxy-2-propyl)valiolamine, a tablet comprising about 5 to 60 mg (preferably about 5 to 30 mg) of N-[N-[(S)-1-ethoxycarbonyl-3-phenylpropyl]-L-alanyl]-N-(indan-2-yl)glycine
20 hydrochloride or a tablet comprising about 0.01 to 1 mg (preferably about 0.01 to 0.5 mg) of (+)-3R,5S-erythro-(E)-7-[4-(4-fluorophenyl)-2,6-diisopropyl-5-methoxymethylpyridin-3-yl]-3,5-dihydroxyhept-6-enoate sodium. Each tablet is preferably administered once a
25 day and may be administered to one and same subject simultaneously or with time intervals of 12 hours or less (preferably 6 hours or less).

(2) A tablet comprising about 1 to 50 mg (preferably about 1 to 35 mg) of 2-ethoxy-1-[[2'-(1H-tetrazol-5-yl)biphenyl-4-yl]methyl]-1H-benzimidazole-7-carboxylic
30 acid is orally administered to one and same subject in the form of combination use with a tablet comprising about 0.1 to 45 mg (preferably about 2 to 30 mg) of 5-[4-[2-(5-ethyl-2-pyridyl)ethoxy]benzyl]-2,4-thiazolidinedione, a tablet comprising about 1 to 20 mg
35 (preferably about 1 to 15 mg) of (R)-(+)-5-[3-[4-[2-(2-

furyl)-5-methyl-4-oxazolylmethoxy]-3-methoxyphenyl]-propyl]-2,4-oxazolidinedione, a tablet comprising about 0.1 to 0.5 mg (preferably about 0.1 to 0.3 mg) of N-(1,3-dihydroxy-2-propyl)valiolamine, a tablet

5 comprising about 5 to 60 mg (preferably about 5 to 30 mg) of N-[N-[(S)-1-ethoxycarbonyl-3-phenylpropyl]-L-alanyl]-N-(indan-2-yl)glycine hydrochloride or a tablet comprising about 0.01 to 0.1 mg (preferably about 0.01 to 0.5 mg) of (+)-3R,5S-erythro-(E)-7-[4-(4-

10 fluorophenyl)-2,6-diisopropyl-5-methoxymethylpyridin-3-yl]-3,5-dihydroxyhept-6-enoate sodium. Each tablet is preferably administered once a day and may be administered to one and same subject simultaneously or with time intervals of 12 hours or less (preferably 6

15 hours or less).

(3) A tablet comprising about 1 to 50 mg (preferably about 1 to 35 mg) of 2-ethoxy-1-[[2'-(2,5-dihydro-5-oxo-1,2,4-oxadiazol-3-yl)biphenyl-4-yl]methyl]-1H-benzimidazole-7-carboxylic acid is orally administered

20 to one and same subject in the form of combination use with a tablet comprising about 0.1 to 45 mg (preferably about 2 to 30 mg) of 5-[4-[2-(5-ethyl-2-pyridyl)ethoxy]benzyl]-2,4-thiazolidinedione, a tablet comprising about 1 to 20 mg (preferably about 1 to 15

25 mg) of (R)-(+)-5-[3-[4-[2-(2-furyl)-5-methyl-4-oxazolylmethoxy]-3-methoxyphenyl]propyl]-2,4-oxazolidinedione, a tablet comprising about 0.1 to 0.5 mg (preferably about 0.1 to 0.3 mg) of N-(1,3-dihydroxy-2-propyl)valiolamine, a tablet comprising

30 about 5 to 60 mg (preferably about 5 to 30 mg) of N-[N-[(S)-1-ethoxycarbonyl-3-phenylpropyl]-L-alanyl]-N-(indan-2-yl)glycine hydrochloride or a tablet comprising about 0.01 to 1 mg (preferably about 0.01 to 0.5 mg) of (+)-3R,5S-erythro-(E)-7-[4-(4-fluorophenyl)-

35 2,6-diisopropyl-5-methoxymethylpyridin-3-yl]-3,5-dihydroxyhept-6-enate sodium. Each tablet is

preferably administered once a day and may be administered to one and same subject simultaneously or with time intervals of 12 hours or less (preferably 6 hours or less).

5 BEST MODE FOR CARRYING OUT THE INVENTION

By the following formulation examples, the present invention will be illustrated in more detail, and they should not be construed as limiting the invention thereto.

10 Examples

Formulation Examples

The pharmaceutical composition (especially the prophylactic or therapeutic agents of angiotensin II-mediated diseases, preferably therapeutic agent for
15 arteriosclerosis of human adult) referred to in this invention, formulated by combination of a compound having the angiotensin II antagonistic activity or a salt thereof with at least one species of a compound having the activity of increasing insulin-sensitivity,
20 a compound having the activity of improving postprandial hyperglycemia in diabetes mellitus, an indane derivative having the activity of inhibiting angiotensin converting enzyme, a pyridine derivative having the activity of inhibiting HMG-Co A reductase or
25 salts thereof can be prepared by, for example, the following prescriptions.

1. Capsules

	(1) 2-ethoxy-1-[[2'-(1H-tetrazol-5-yl)biphenyl-4-yl]-methyl]-1H-benzimidazole-7-carboxylic acid	1 mg
30	(2) 5-[4-[2-(5-ethyl-2-pyridyl)ethoxy]benzyl]-2,4-thiazolidinedione	30 mg
	(3) lactose	69 mg
	(4) microcrystalline cellulose	70 mg
35	(5) magnesium stearate	10 mg
one capsule 180 mg		

(1), (2), (3), (4) and 1/2 of (5) were mixed and then granulated. To the granules was added the remainder of (5), and the whole was filled into a gelatin capsule.

2. Tablets

- 5 (1) 2-ethoxy-1-[[2'-(1H-tetrazol-5-yl)biphenyl-4-yl]-methyl]-1H-benzimidazole-7-carboxylic acid

1 mg

- (2) 5-[4-[2-(5-ethyl-2-pyridyl)ethoxy]benzyl-2,4-thiazolidinedione

30 mg

- 10 (3) lactose 66.4 mg

- (4) corn starch 20 mg

- (5) polyethylene glycol 2.6 mg

- (6) hydroxypropyl cellulose 4 mg

- (7) carmellose calcium 5.6 mg

- 15 (8) magnesium stearate 0.4 mg

one tablet 130 mg

(1), (2), (3), (4), (5), 2/3 of (6), 2/3 of (7) and 1/2 of (8) were mixed and then granulated. To the granules were added the remainders of (6), (7) and (8), followed by subjecting the mixture to compression molding.

3. Injections

- (1) 2-ethoxy-1-[[2'-(1H-tetrazol-5-yl)biphenyl-4-yl]-methyl]-1H-benzimidazole-7-carboxylic acid

25 1 mg

- (2) 5-[4-[2-(5-ethyl-2-pyridyl)ethoxy]benzyl]-

2,4-thiazolidinedione 30 mg

- (3) inositol 79 mg

- (4) benzyl alcohol 20 mg

30 one ampoule 130 mg

(1), (2), (3) and (4) were dissolved in distilled water for injection to make the whole volume 2 ml, which was filled into an ampoule. The whole process was conducted under sterile conditions.

4. Capsules

- (1) 2-ethoxy-1-[[2'-(1H-tetrazol-5-yl)biphenyl-4-yl]-

methyl]benzimidazole-7-carboxylic acid

1 mg

(2) (R)-(+)-5-[3-[4-[2-(2-furyl)-5-methyl-4-oxazolylmethoxy]-3-methoxyphenyl]propyl]-2,4-

5 oxazolidinedione 10 mg

(3) lactose 89 mg

(4) microcrystalline cellulose 70 mg

(5) magnesium stearate 10 mg

one capsule 180 mg

10 (1), (2), (3), (4) and 1/2 of (5) were mixed and then granulated. To the granules was added the remainder of (5), and the whole was filled into a gelatin capsule.

5. Tablets

15 (1) 2-ethoxy-1-[[2'-(1H-tetrazol-5-yl)biphenyl-4-yl]-methyl]-1H-benzimidazole-7-carboxylic acid 1 mg

(2) (R)-(+)-5-[3-[4-[2-(2-furyl)-5-methyl-4-oxazolylmethoxy]-3-methoxyphenyl]propyl]-2,4-oxazolidinedione 10 mg

20 (3) lactose 86.4 mg

(4) corn starch 20 mg

(5) polyethylene glycol 2.6 mg

(6) hydroxypropyl cellulose 4 mg

(7) carmellose calcium 5.6 mg

25 (8) magnesium stearate 0.4 mg

one tablet 130 mg

(1), (2), (3), (4), (5), 2/3 of (6), 2/3 of (7) and 1/2 of (8) were mixed and then granulated. To the granules were added the remainders of (6), (7) and (8), followed by subjecting the mixture to compression molding.

6. Injections

(1) 2-ethoxy-1-[[2'-(1H-tetrazol-5-yl)biphenyl-4-yl]methyl]-1H-benzimidazole-7-carboxylic acid 1 mg

35

(2) (R)-(+)-5-[3-[4-[2-(2-furyl)-5-methyl-4-

	oxazolylmethoxy]-3-methoxyphenyl]propyl]-	
	2,4-oxazolidinedione	10 mg
	(3) inositol	99 mg
	(4) benzyl alcohol	20 mg
5	one ampoule	130 mg

(1), (2), (3) and (4) were dissolved in distilled water for injection to make the whole volume 2 ml, which is filled into an ampoule. The whole process was conducted under sterile conditions.

10 7. Capsules

	(1) 2-ethoxy-1-[[2'-(1H-tetrazol-5-yl)biphenyl-4-yl]-methyl]-1H-benzimidazole-7-carboxylic acid	
		1 mg
	(2) N-(1,3-dihydroxy-2-propyl)valiolamine	0.2 mg
15	(3) lactose	98.8 mg
	(4) microcrystalline cellulose	70 mg
	(5) magnesium stearate	10 mg
	one capsule	180 mg

(1), (2), (3), (4) and 1/2 of (5) were mixed and then granulated. To the granules was added the remainder of (5), and the whole was filled into a gelatin capsule.

20 8. Tablets

	(1) 2-ethoxy-1-[[2'-(1H-tetrazol-5-yl)biphenyl-4-yl]-methyl]-1H-benzimidazole-7-carboxylic acid	
25		1 mg
	(2) N-(1,3-dihydroxy-2-propyl)valiolamine	0.2 mg
	(3) lactose	96.2 mg
	(4) corn starch	20 mg
30	(5) polyethylene glycol	2.6 mg
	(6) hydroxypropyl cellulose	4 mg
	(7) carmellose calcium	5.6 mg
	(8) magnesium stearate	0.4 mg

one tablet 130 mg

35 (1), (2), (3), (4), (5), 2/3 of (6), 2/3 of (7) and 1/2 of (8) were mixed and then granulated. To the

granules were added the remainders of (6), (7) and (8), followed by subjecting the mixture to compression molding.

9. Injections

- 5 (1) 2-ethoxy-1-[[2'-(1H-tetrazol-5-yl)biphenyl-4-yl]-methyl]-1H-benzimidazole-7-carboxylic acid 1 mg
- (2) N-(1,3-dihydroxy-2-propyl)valiolamine 0.2 mg
- (3) inositol 108.8 mg
- 10 (4) benzyl alcohol 20 mg
- one ampoule 130 mg

(1), (2), (3) and (4) were dissolved in distilled water for injection to make the whole volume 2 ml, which was filled into an ampoule. The whole process was conducted under sterile conditions.

10. Capsules

- 15 (1) 2-ethoxy-1-[[2'-(1H-tetrazol-5-yl)biphenyl-4-yl]-methyl]benzimidazole-7-carboxylic acid 1 mg
- (2) N-[N-[(S)-1-ethoxycarbonyl-3-phenylpropyl]-L-alanyl]-N-(indan-2-yl)glycine hydrochloride 10 mg
- 20 (3) lactose 89 mg
- (4) microcrystalline cellulose 70 mg
- (5) magnesium stearate 10 mg
- 25 one capsule 180 mg

(1), (2), (3), (4) and 1/2 of (5) were mixed and then granulated. To the granules was added the remainder of (5), and the whole was filled into a gelatin capsule.

30 11. Tablets

- (1) 2-ethoxy-1-[[2'-(1H-tetrazol-5-yl)biphenyl-4-yl]-methyl]1H-benzimidazole-7-carboxylic acid 1 mg
- (2) N-[N-[(S)-1-ethoxycarbonyl-3-phenylpropyl]-L-alanyl]-N-(indan-2-yl)glycine hydrochloride 10 mg
- 35 (3) lactose 86.4 mg

	(4) corn starch	20 mg
	(5) polyethylene glycol	2.6 mg
	(6) hydroxypropyl cellulose	4 mg
	(7) carmellose calcium	5.6 mg
5	(8) magnesium stearate	0.4 mg

one tablet 130 mg

(1), (2), (3), (4), (5), 2/3 of (6), 2/3 of (7) and 1/2 of (8) were mixed and then granulated. To the granules were added the remainders of (6), (7) and (8), followed by subjecting the mixture to compression molding.

12. Injections

	(1) 2-ethoxy-1-[[2'-(1H-tetrazol-5-yl)biphenyl-4-yl]-methyl]-1H-benzimidazole-7-carboxylic acid	1 mg
15	(2) N-[N-[(S)-1-ethoxycarbonyl-3-phenylpropyl]-L-alanyl]-N-(indan-2-yl)glycine hydrochloride	
		10 mg
	(3) inositol	99 mg
	(4) benzyl alcohol	20 mg

20 one ampoule 130 mg

(1), (2), (3) and (4) were dissolved in distilled water for injection to make the whole volume 2 ml, which was filled into an ampoule. The whole process was conducted under sterile conditions.

25 13. Capsules

	(1) 2-ethoxy-1-[[2'-(1H-tetrazol-5-yl)biphenyl-4-yl]-methyl]-1H-benzimidazole-7-carboxylic acid	1 mg
30	(2) (+)-3R,5S-erythro-(E)-7-[4-(4-fluorophenyl)-2,6-diisopropyl-5-methoxymethylpyridin-3-yl]-3,5-dihydroxyhept-6-enoate sodium	0.15 mg
	(3) lactose	98.85 mg
	(4) microcrystalline cellulose	70 mg
	(5) magnesium stearate	10 mg

35 one capsule 180 mg

(1), (2), (3), (4) and 1/2 of (5) were mixed and

then granulated. To the granules was added the remainder of (5), and the whole was filled into a gelatin capsule.

14. Tablets

- | | | |
|----|--|----------|
| 5 | (1) 2-ethoxy-1-[[2'-(1H-tetrazol-5-yl)biphenyl-4-yl]-methyl]-1H-benzimidazol-7-carboxylic acid | 1 mg |
| 10 | (2) (+)-3R,5S-erythro-(E)-7-[4-(4-fluorophenyl)-2,6-diisopropyl-5-methoxymethylpyridin-3-yl]-3,5-dihydroxyhept-6-enoate sodium | 0.15 mg |
| | (3) lactose | 96.25 mg |
| | (4) corn starch | 20 mg |
| | (5) polyethylene glycol | 2.6 mg |
| | (6) hydroxypropyl cellulose | 4 mg |
| 15 | (7) carmellose calcium | 5.6 mg |
| | (8) magnesium stearate | 0.4 mg |
| | one tablet | 130 mg |
- (1), (2), (3), (4), (5), 2/3 of (6), 2/3 of (7) and 1/2 of (8) were mixed and then granulated. To the granules were added the remainders of (6), (7) and (8), followed by subjecting the mixture to compression molding.

15. Injections

- | | | |
|----|--|-----------|
| 25 | (1) 2-ethoxy-1-[[2'-(1H-tetrazol-5-yl)biphenyl-4-yl]-methyl]-1H-benzimidazole-7-carboxylic acid | 1 mg |
| | (2) (+)-3R,5S-erythro-(E)-7-[4-(4-fluorophenyl)-2,6-diisopropyl-5-methoxymethylpyridin-3-yl]-3,5-dihydroxyhept-6-enoate sodium | 0.15 mg |
| 30 | (3) inositol | 108.85 mg |
| | (4) benzyl alcohol | 20 mg |
| | one ampoule | 130 mg |
- (1), (2), (3) and (4) were dissolved in distilled water for injection to make the whole volume 2 ml, which is filled into an ampoule. The whole process was conducted under sterile conditions.

16. Capsules

	(1) (\pm)-1-(cyclohexyloxycarbonyloxy)ethyl 2-ethoxy-1-[[2'-(1H-tetrazol-5-yl)biphenyl-4-yl]- methyl]-1H-benzimidazole-7-carboxylate	1 mg
5	(2) 5-[4-[2-(5-ethyl-2-pyridyl)ethoxy]benzyl]- 2,4-thiazolidinedione	30 mg
	(3) lactose	69 mg
	(4) microcrystalline cellulose	70 mg
	(5) magnesium stearate	10 mg
10		one capsule 180 mg

(1), (2), (3), (4) and 1/2 of (5) were mixed and then granulated. To the granules was added the remainder of (5), and the whole was filled into a gelatin capsule.

15 17. Tablets

	(1) (\pm)-1-(cyclohexyloxycarbonyloxy)ethyl 2-ethoxy-1- [[2'-(1H-tetrazol-5-yl)biphenyl-4-yl]methyl]-1H- benzimidazole-7-carboxylate	1 mg
20	(2) 5-[4-[2-(5-ethyl-2-pyridyl)ethoxy]benzyl]- 2,4-thiazolidinedione	30 mg
	(3) lactose	66.4 mg
	(4) corn starch	20 mg
	(5) polyethylene glycol	2.6 mg
	(6) hydroxypropyl cellulose	4 mg
25	(7) carmellose calcium	5.6 mg
	(8) magnesium stearate	0.4 mg

one tablet 130 mg

(1), (2), (3), (4), (5), 2/3 of (6), 2/3 of (7) and 1/2 of (8) were mixed and then granulated. To the granules were added the remainders of (6), (7) and (8), followed by subjecting the mixture to compression molding.

18. Capsules

35	(1) (\pm)-1-(cyclohexyloxycarbonyloxy)ethyl 2-ethoxy-1- [[2'-(1H-tetrazol-5-yl)biphenyl-4-yl]methyl]-1H- benzimidazole-7-carboxylate	1 mg
----	--	------

- (2) (R)-(+)-5-[3-[4-[2-(2-furyl)-5-methyl-4-oxazolylmethoxy]-3-methoxyphenyl]propyl-2,4-oxazolidinedione 10 mg
- (3) lactose 89 mg
- 5 (4) microcrystalline cellulose 70 mg
- (5) magnesium stearate 10 mg
- one capsule 180 mg

(1), (2), (3), (4) and 1/2 of (5) were mixed and then granulated. To the granules was added the remainder of (5), and the whole was filled into a gelatin capsule.

19. Tablets

- (1) (±)-1-(cyclohexyloxycarbonyloxy)ethyl 2-ethoxy-1-[[2'-(1H-tetrazol-5-yl)biphenyl-4-yl]methyl]-1H-benzimidazole-7-carboxylate 1 mg
- 15 (2) (R)-(+)-5-[3-[4-[2-(2-furyl)-5-methyl-4-oxazolylmethoxy]-3-methoxyphenyl]propyl]-2,4-oxazolidinedione 10 mg
- (3) lactose 86.4 mg
- 20 (4) corn starch 20 mg
- (5) polyethylene glycol 2.6 mg
- (6) hydroxypropyl cellulose 4 mg
- (7) carmellose calcium 5.6 mg
- (8) magnesium stearate 0.4 mg

25 one tablet 130 mg

(1), (2), (3), (4), (5), 2/3 of (6), 2/3 of (7) and 1/2 of (8) were mixed and then granulated. To the granules were added the remainders of (6), (7) and (8), followed by subjecting the mixture to compression molding.

20. Capsules

- (1) (±)-1-(cyclohexyloxycarbonyloxy)ethyl 2-ethoxy-1-[[2'-(1H-tetrazol-5-yl)biphenyl-4-yl]methyl]-1H-benzimidazole-7-carboxylate 1 mg
- 35 (2) N-(1,3-dihydroxy-2-propyl)valiolamine 0.2 mg
- (3) lactose 98.8 mg

- (4) microcrystalline cellulose 70 mg
 (5) magnesium stearate 10 mg

one capsule 180 mg

(1), (2), (3), (4) and 1/2 of (5) were mixed and
 5 then granulated. To the granules was added the
 remainder of (5), and the whole was filled into a
 gelatin capsule.

21. Tablets

- (1) (\pm)-1-(cyclohexyloxycarbonyloxy)ethyl 2-ethoxy-1-
 10 [[2'-(1H-tetrazol-5-yl)biphenyl-4-yl]methyl]-1H-
 benzimidazole-7-carboxylate 1 mg
 (2) N-(1,3-dihydroxy-2-propyl)valiolamine 0.2 mg
 (3) lactose 96.2 mg
 (4) corn starch 20 mg
 15 (5) polyethylene glycol 2.6 mg
 (6) hydroxypropyl cellulose 4 mg
 (7) carmellose calcium 5.6 mg
 (8) magnesium stearate 0.4 mg

one tablet 130 mg

(1), (2), (3), (4), (5), 2/3 of (6), 2/3 of (7)
 20 and 1/2 of (8) were mixed and then granulated. To the
 granules were added the remainders of (6), (7) and (8),
 followed by subjecting the mixture to compressive
 molding.

22. Capsules

- (1) (\pm)-1-(cyclohexyloxycarbonyloxy)ethyl 2-ethoxy-1-
 [[2'-(1H-tetrazol-5-yl)biphenyl-4-yl]methyl]-1H-
 benzimidazole-7-carboxylate 1 mg
 (2) N-[N-[(S)-1-ethoxycarbonyl-3-phenylpropyl]-L-
 30 alanyl-N-(indan-2-yl)glycine hydrochloride 10 mg
 (3) lactose 89 mg
 (4) microcrystalline cellulose 70 mg
 (5) magnesium stearate 10 mg

35 one capsule 180 mg

(1), (2), (3), (4) and 1/2 of (5) were mixed and

then granulated. To the granules was added the remainder of (5), and the whole was filled into a gelatin capsule.

23. Tablets

5	(1) (\pm)-1-(cyclohexyloxycarbonyloxy)ethyl 2-ethoxy-1- [[2'-(1H-tetrazol-5-yl)biphenyl-4-yl]methyl]- 1H-benzimidazole-7-carboxylate	1 mg
	(2) N-[N-[(S)-1-ethoxycarbonyl-3-phenylpropyl]-L- alanyl]-N-(indan-2-yl)glycine hydrochloride	
10		10 mg
	(3) lactose	86.4 mg
	(4) corn starch	20 mg
	(5) polyethylene glycol	2.6 mg
	(6) hydroxypropyl cellulose	4 mg
15	(7) carmellose calcium	5.6 mg
	(8) magnesium stearate	0.4 mg
		one tablet 130 mg

(1), (2), (3), (4), (5), 2/3 of (6), 2/3 of (7) and 1/2 of (8) were mixed and then granulated. To the granules were added the remainders of (6), (7) and (8), followed by subjecting the mixture to compression molding.

24. Capsules

25	(1) (\pm)-1-(cyclohexyloxycarbonyloxy)ethyl 2-ethoxy-1- [[2'-(1H-tetrazol-5-yl)biphenyl-4-yl]methyl]-1H- benzimidazole-7-carboxylate	1 mg
	(2) (+)-3R,5S-erythro-(E)-7-[4-(4-fluorophenyl)- 2,6-diisopropyl-5-methoxymethylpyridin-3-yl]- 3,5-dihydroxyhept-6-enoate sodium	0.15 mg
30	(3) lactose	98.85 mg
	(4) microcrystalline cellulose	70 mg
	(5) magnesium stearate	10 mg
		one capsule 180 mg

(1), (2), (3), (4) and 1/2 of (5) were mixed and then granulated. To the granules was added the remainder of (5), and the whole was filled into a

gelatin capsule.

25. Tablets

- (1) (\pm)-1-(cyclohexyloxycarbonyloxy)ethyl 2-ethoxy-1-
 [[2'-(1H-tetrazol-5-yl)biphenyl-4-yl]methyl]-1H-
 5 benzimidazole-7-carboxylate 1 mg
- (2) (+)-3R,5S-erythro-(E)-7-[4-(4-fluorophenyl)-
 2,6-diisopropyl-5-methoxymethylpyridin-3-yl]-
 3,5-dihydroxyhept-6-enoate sodium 0.15 mg
- (3) lactose 96.25 mg
- 10 (4) corn starch 20 mg
- (5) polyethylene glycol 2.6 mg
- (6) hydroxypropyl cellulose 4 mg
- (7) carmellose calcium 5.6 mg
- (8) magnesium stearate 0.4 mg

15 one tablet 130 mg

(1), (2), (3), (4), (5), 2/3 of (6), 2/3 of (7)
 and 1/2 of (8) were mixed and then granulated. To the
 granules were added the remainders of (6), (7) and (8),
 followed by subjecting the mixture to compression
 20 molding.

26. Capsules

- (1) 2-ethoxy-1-[[2'-(2,5-dihydro-5-oxo-1,2,4-
 oxadiazol-3-yl)biphenyl-4-yl]methyl]-1H-
 benzimidazole-7-carboxylic acid 1 mg
- 25 (2) 5-[4-[2-(5-ethyl-2-pyridyl)ethoxy]benzyl]-
 2,4-thiazolidinedione 30 mg
- (3) lactose 69 mg
- (4) microcrystalline cellulose 70 mg
- (5) magnesium stearate 10 mg

30 one capsule 180 mg

(1), (2), (3), (4) and 1/2 of (5) were mixed and
 then granulated. To the granules was added the
 remainder of (5), and the whole was filled into a
 gelatin capsule.

35 27. Tablets

- (1) 2-ethoxy-1-[[2'-(2,5-dihydro-5-oxo-1,2,4-oxadiazol-

	3-yl)biphenyl-4-yl)methyl]-1H-benzimidazole-7-carboxylic acid	1 mg
	(2) 5-[4-[2-(5-ethyl-2-pyridyl)ethoxy]benzyl]-2,4-thiazolidinedione	30 mg
5	(3) lactose	66.4 mg
	(4) corn starch	20 mg
	(5) polyethylene glycol	2.6 mg
	(6) hydroxypropyl cellulose	4 mg
	(7) carmellose calcium	5.6 mg
10	(8) magnesium stearate	0.4 mg
	one tablet	130 mg

(1), (2), (3), (4), (5), 2/3 of (6), 2/3 of (7) and 1/2 of (8) were mixed and then granulated. To the granules were added the remainders of (6), (7) and (8), followed by subjecting the mixture to compression molding.

28. Capsules

	(1) 2-ethoxy-1-[[2'-(2,5-dihydro-5-oxo-1,2,4-oxadiazol-3-yl)biphenyl-4-yl)methyl]-1H-benzimidazole-7-carboxylic acid	1 mg
20	(2) (R)-(+)-5-[3-[4-[2-furyl)-5-methyl-4-oxazolylmethoxy]-3-methoxyphenyl]propyl]-2,4-oxazolidinedione	10 mg
	(3) lactose	89 mg
25	(4) microcrystalline cellulose	70 mg
	(5) magnesium stearate	10 mg
	one capsule	180 mg

(1), (2), (3), (4) and 1/2 of (5) were mixed and then granulated. To the granules was added the remainder of (5), and the whole was filled into a gelatin capsule.

29. Tablets

	(1) 2-ethoxy-1-[[2'-(2,5-dihydro-5-oxo-1,2,4-oxadiazol-3-yl)biphenyl-4-yl)methyl]-1H-benzimidazole-7-carboxylic acid	1 mg
35	(2) (R)-(+)-5-[3-[4-[2-(2-furyl)-5-methyl-4-	

	oxazolylmethoxy]-3-methoxyphenyl]propyl]-	
	2,4-oxazolidinedione	10 mg
	(3) lactose	86.4 mg
	(4) corn starch	20 mg
5	(5) polyethylene glycol	2.6 mg
	(6) hydroxypropyl cellulose	4 mg
	(7) carmellose calcium	5.6 mg
	(8) magnesium stearate	0.45 mg
	one tablet	130 mg
10	(1), (2), (3), (4), (5) 2/3 of (6), 2/3 of (7)	
	and 1/2 of (8) were mixed and then granulated. To the	
	granules were added the remainders of (6), (7) and (8),	
	followed by subjecting the mixture to compression	
	molding.	
15	30. Capsules	
	(1) 2-ethoxy-1-[[2'-(2,5-dihydro-5-oxo-1,2,4-oxadiazol-	
	3-yl]biphenyl-4-yl)methyl]-1H-benzimidazole-7-	
	carboxylic acid	1 mg
	(2) N-(1,3-dihydroxy-2-propyl)valiolamine	0.2 mg
20	(3) lactose	98.8 mg
	(4) microcrystalline cellulose	70 mg
	(5) magnesium stearate	10 mg
	one capsule	180 mg
	(1), (2), (3), (4) and 1/2 of (5) were mixed and	
25	then granulated. To the granules was added the	
	remainder of (5), and the whole was filled into a	
	gelatin capsule,	
	31. Tablets	
	(1) 2-ethoxy-1-[[2'-(2,5-dihydro-5-oxo-1,2,4-oxadiazol-	
30	3-yl)biphenyl-4-yl)methyl]-1H-benzimidazole-7-	
	carboxylic acid	1 mg
	(2) N-(1,3-dihydroxy-2-propyl)valiolamine	0.2 mg
	(3) lactose	96.2 mg
	(4) corn starch	20 mg
35	(5) polyethylene glycol	2.6 mg
	(6) hydroxypropyl cellulose	4 mg

(7) carmellose calcium	5.6 mg
(8) magnesium stearate	0.4 mg

one tablet 130 mg

(1), (2), (3), (4), (5), 2/3 of (6), 2/3 of (7) and 1/2 of (8) were mixed and then granulated. To the granules were added the remainders of (6), (7) and (8), followed by subjecting the mixture to compression molding.

32. Capsules

10	(1) 2-ethoxy-1-[[2'-(2,5-dihydro-5-oxo-1,2,4-oxadiazol-3-yl)biphenyl-4-yl]methyl]-1H-benzimidazole-7-carboxylic acid	1 mg
	(2) N-[N-[(S)-1-ethoxycarbonyl-3-phenylpropyl]-L-alanyl]-N-(indan-2-yl)glycine hydrochloride	

15		10 mg
	(3) lactose	89 mg
	(4) microcrystalline cellulose	70 mg
	(5) magnesium stearate	10 mg

one capsule 180 mg

20 (1), (2), (3), (4) and 1/2 of (5) were mixed and then granulated. To the granules was added the remainder of (5), and the whole was filled into a gelatin capsule.

33. Tablets

25	(1) 2-ethoxy-1-[[2'-(2,5-dihydro-5-oxo-1,2,4-oxadiazol-3-yl)biphenyl-4-yl]methyl]-1H-benzimidazole-7-carboxylic acid	1 mg
	(2) N-[N-[(S)-1-ethoxycarbonyl-3-phenylpropyl]-L-alanyl]-N-(indan-2-yl)glycine hydrochloride	
30		10 mg
	(3) lactose	86.4 mg
	(4) corn starch	20 mg
	(5) polyethylene glycol	2.6 mg
	(6) hydroxypropyl cellulose	4 mg
35	(7) carmellose calcium	5.6 mg
	(8) magnesium stearate	0.4 mg

one tablet 130 mg

(1), (2), (3), (4), (5), 2/3 of (6), 2/3 of (7) and 1/2 of (8) were mixed and then granulated. To the granules were added the remainders of (6), (7) and (8), followed by subjecting the mixture to compression molding.

34. Capsules

- | | | |
|----|--|----------|
| 10 | (1) 2-ethoxy-1-[[2'-(2,5-dihydro-5-oxo-1,2,4-oxadiazol-3-yl)biphenyl-4-yl]methyl]-1H-benzimidazole-7-carboxylic acid | 1 mg |
| | (2) (+)-3R,5S-erythro-(E)-7-[4-(4-fluorophenyl)-2,6-diisopropyl-5-methoxymethylpyridin-3-yl]-3,5-dihydroxyhept-6-enoate sodium | 0.15 mg |
| | (3) lactose | 98.85 mg |
| 15 | (4) microcrystalline cellulose | 70 mg |
| | (5) magnesium stearate | 10 mg |

one capsule 180 mg

- (1), (2), (3), (4) and 1/2 of (5) were mixed and then granulated. To the granules was added the remainder of (5), and the whole was filled into a gelatin capsule.

35. Tablets

- | | | |
|----|--|----------|
| 25 | (1) 2-ethoxy-1-[[2'-(2,5-dihydro-5-oxo-1,2,4-oxadiazole-3-yl)biphenyl-4-yl]methyl]-1H-benzimidazol-7-carboxylic acid | 1 mg |
| | (2) (+)-3R,5S-erythro-(E)-7-[4-(4-fluorophenyl)-2,6-diisopropyl-5-methoxymethylpyridin-3-yl]-3,5-dihydroxyhept-6-enoate sodium | 0.15 mg |
| | (3) lactose | 96.25 mg |
| 30 | (4) corn starch | 20 mg |
| | (5) polyethylene glycol | 2.6 mg |
| | (6) hydroxypropyl cellulose | 4 mg |
| | (7) carmellose calcium | 5.6 mg |
| | (8) magnesium stearate | 0.4 mg |

one tablet 130 mg

(1), (2), (3), (4), (5), 2/3 of (6), 2/3 of (7)

and 1/2 of (8) were mixed and then granulated. To the granules were added the remainders of (6), (7) and (8), followed by subjecting the mixture to compression molding.

5 INDUSTRIAL APPLICABILITY

The pharmaceutical composition of this invention formulated by combination of an angiotensin II antagonistic compound or a salt thereof with at least one species of a compound having the activity of increasing insulin-sensitivity, a compound having the activity of improving postprandial hyperglycemia in diabetes mellitus, an indane derivative having the activity of inhibiting angiotensin converting enzyme, a pyridine derivative having the activity of inhibiting HMG-Co A reductase or salts thereof serves to decrease remarkably the dosages of the individual effective components, and, as a result, suppresses undesirable side effects observed in the case of administering the respective compounds singly, and can be advantageously used as a prophylactic or therapeutic agent of angiotensin II-mediated diseases, especially arteriosclerosis or arteriosclerosis having hypertension as a complication.

CLAIMS

1. A pharmaceutical composition comprising a compound having angiotensin II antagonistic activity or a salt thereof in combination with at least one species selected from the group consisting of a compound having the activity of increasing insulin-sensitivity, a compound having the activity of lowering postprandial hyperglycemia in diabetes mellitus, an indane derivative having the activity of inhibiting angiotensin converting enzyme, a pyridine derivative having the activity of inhibiting HMG-Co A reductase and their salts.
2. The composition as claimed in Claim 1, which is a preventing or treating agent of angiotensin II-mediated diseases.
3. The composition as claimed in Claim 2, which is directed to the prevention or treatment of circulatory diseases.
4. The composition as claimed in Claim 2, which is directed to the prevention or treatment of hypertension, cardiac insufficiency, cerebral apoplexy, ischemic peripheral circulation disturbances, myocardial ischemia, venous insufficiency, progressive cardiac insufficiency after myocardial infarction, diabetic nephropathy, nephritis, glomerulonephritis, arteriosclerosis, angiohypertrophy, vascular hypertrophy or obstruction after percutaneous transluminal coronary angioplasty, vascular reobstruction after bypass surgery, hyperaldosteronism, glomerulosclerosis, renal insufficiency, glaucoma, ocular hypertension, hyperlipemia, myocardial infarction, angina pectoris, aneurysm, coronary arteriosclerosis, cerebral arteriosclerosis, peripheral arteriosclerosis, thrombosis, diseases of central nervous system, Alzheimer's disease, deficiency of memory, depression, amnesia, senile dementia, sensory

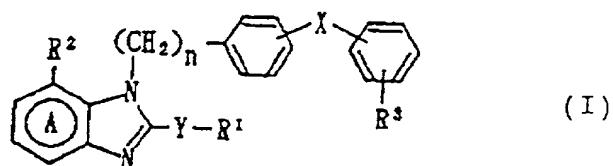
disturbances, multiple system organ failure or scleroderma, or to the prevention or amelioration of anxiety neurosis, catatonia, indisposition or dyspeptic symptoms.

5. The composition as claimed in Claim 2, which is directed to the prevention or treatment of complications of hypertension.

6. The composition as claimed in Claim 2, which is directed to the prevention or treatment of arterio-sclerosis.

7. The composition as claimed in Claim 5, which is directed to the prevention or treatment of arterio-sclerosis.

8. The composition as claimed in Claim 1, wherein the compound having angiotensin II antagonistic activity is a compound of the formula:



wherein R^1 stands for H or an optionally substituted hydrocarbon residue; R^2 stands for an optionally esterified carboxyl group; R^3 stands for a group capable of forming anion or a group convertible thereto; X shows that phenylene group and phenyl group are bonded directly or through a spacer having a chain length of 1 to 2 atoms; n denotes 1 or 2; the ring A is a benzene ring optionally having further substituents other than the group shown by R^2 ; and Y stands for a bond, -O-, -S(O)m- (m denotes 0, 1 or 2) or -N(R^4)- (R^4 stands for H or an optionally substituted alkyl group).

9. The composition as claimed in Claim 1, wherein the compound having angiotensin II antagonistic activity is (+)-1-(cyclohexyloxycarbonyloxy)ethyl 2-ethoxy-1-[[2'-(1H-tetrazol-5-yl)biphenyl-4-yl]methyl]-

14. The composition as claimed in Claim 1, wherein the compound having angiotensin II antagonistic activity is (\pm) -1-(cyclohexyloxycarbonyloxy)ethyl 2-ethoxy-1-[[2'-(1H-tetrazol-5-yl)biphenyl-4-yl]methyl]-1H-benzimidazole-7-carboxylate, 2-ethoxy-1-[[2'-(1H-tetrazol-5-yl)biphenyl-4-yl]methyl]-1H-benzimidazole-7-carboxylic acid or 2-ethoxy-1-[[2'-(2,5-dihydro-5-oxo-1,2,4-oxadiazol-3-yl)biphenyl-4-yl]methyl]-1H-benzimidazole-7-carboxylic acid;

the compound having the activity of increasing insulin-sensitivity is 5-[4-[2-(5-ethyl-2-pyridyl)ethoxy]-

benzyl]-2,4-thiazolidinedione or (R)-(+)-5-[3-[4-[2-(2-furyl)-5-methyl-4-oxazolylmethoxy]-3-methoxyphenyl]-propyl]-2,4-oxazolidinedione;

the compound having the activity of improving postprandial hyperglycemia in diabetes mellitus is N-(1,3-dihydroxy-2-propyl)valiolamine;

the indane derivative having the activity of inhibiting angiotensin converting enzyme is N-[N-[(S)-1-ethoxycarbonyl-3-phenylpropyl]-L-alanyl]-N-(indan-2-yl)-glycine; and

the pyridine derivative having the activity of inhibiting HMG-Co A reductase is (+)-3R,5S-erythro-(E)-7-[4-(4-fluorophenyl)-2,6-diisopropyl-5-methoxymethylpyridin-3-yl]-3,5-dihydroxyhept-6-enoic acid.

15. The composition as claimed in Claim 1 comprising the compound having angiotensin II antagonistic activity or a salt thereof in combination with the compound having the activity of increasing insulin-sensitivity or a salt thereof.

16. The composition as claimed in Claim 1 comprising the compound having angiotensin II antagonistic activity or a salt thereof in combination with the compound having the activity of lowering postprandial hyperglycemia in diabetes mellitus or a salt thereof.

17. A pharmaceutical composition for the prevention or treatment of hypertension, arteriosclerosis or hyperlipemia comprising (±)-1-(cyclohexyloxycarbonyloxy)ethyl 2-ethoxy-1-[[2'-(1H-tetrazol-5-yl)biphenyl-4-yl]methyl]-1H-benzimidazole-7-carboxylate or a salt thereof in combination with at least one species selected from the group consisting of 5-[4-[2-(5-ethyl-2-pyridyl)ethoxy]benzyl]-2,4-thiazolidinedione, (R)-(+)-5-[3-[4-[2-(2-furyl)-5-methyl-4-oxazolylmethoxy]-3-methoxyphenyl]propyl]-2,4-oxazolidinedione, N-(1,3-dihydroxy-2-propyl)valiolamine, N-[N-[(S)-1-ethoxycarbonyl-3-phenylpropyl]-L-alanyl]-N-

(indan-2-yl)glycine, (+)-3R,5S-erythro-(E)-7-[4-(4-fluorophenyl)-2,6-diisopropyl-5-methoxymethylpyridin-3-yl]-3,5-dihydroxyhept-6-enoic acid and their salts.

18. A pharmaceutical composition for the prevention or treatment of hypertension, arteriosclerosis or hyperlipemia comprising 2-ethoxy-1-[[2'-(1H-tetrazol-5-yl)biphenyl-4-yl]methyl]-1H-benzimidazole-7-carboxylic acid or a salt thereof in combination with at least one species selected from the group consisting of 5-[4-[2-(5-ethyl-2-pyridyl)ethoxy]benzyl]-2,4-thiazolidinedione, (R)-(+)-5-[3-[4-[2-(2-furyl)-5-methyl-4-oxazolylmethoxy]-3-methoxyphenyl]-propyl]-2,4-oxazolidinedione, N-(1,3-dihydroxy-2-propyl)valiolamine, N-[N-[(S)-1-ethoxycarbonyl-3-phenylpropyl]-L-alanyl]-N-(indan-2-yl)glycine, (+)-3R,5S-erythro-(E)-7-[4-(4-fluorophenyl)-2,6-diisopropyl-5-methoxymethylpyridin-3-yl]-3,5-dihydroxyhept-6-enoic acid and their salts.

19. A pharmaceutical composition for the prevention or treatment of hypertension, arteriosclerosis or hyperlipemia comprising 2-ethoxy-1-[[2'-(2,5-dihydro-5-oxo-1,2,4-oxadiazol-3-yl)biphenyl-4-yl]methyl]-1H-benzimidazole-7-carboxylic acid or a salt thereof in combination with at least one species selected from the group consisting of 5-[4-[2-(5-ethyl-2-pyridyl)ethoxy]benzyl]-2,4-thiazolidinedione, (R)-(+)-5-[3-[4-[2-(2-furyl)-5-methyl-4-oxazolylmethoxy]-3-methoxyphenyl]propyl]-2,4-oxazolidinedione, N-(1,3-dihydroxy-2-propyl)valiolamine, N-[N-[(S)-1-ethoxycarbonyl-3-phenylpropyl]-L-alanyl]-N-(indan-2-yl)glycine, (+)-3R,5S-erythro-(E)-7-[4-(4-fluorophenyl)-2,6-diisopropyl-5-methoxymethylpyridin-3-yl]-3,5-dihydroxyhept-6-enoic acid and their salts.

20. A method for preventing or treating angiotensin II-mediated diseases in a mammal, which comprises administering to said mammal a compound having

angiotensin II antagonistic activity or a salt thereof in combination with at least one species selected from the group consisting of a compound having the activity of increasing insulin-sensitivity, a compound having the activity of lowering postprandial hyperglycemia in diabetes mellitus, an indane derivative having the activity of inhibiting angiotensin converting enzyme, a pyridine derivative having the activity of inhibiting HMG-Co A reductase and their salts.

21. Use of a compound having angiotensin II antagonistic activity or a salt thereof in combination with at least one species selected from the group consisting of a compound having the activity of increasing insulin-sensitivity, a compound having the activity of lowering postprandial hyperglycemia in diabetes mellitus, an indane derivative having the activity of inhibiting angiotensin converting enzyme, a pyridine derivative having the activity of inhibiting HMG-Co A reductase and their salts, for the manufacture of a medicament for preventing or treating angiotensin II-mediated diseases.

ABSTRACT

To provide a pharmaceutical composition which performs a remarkable effect with a relatively decreased dosage, and, with less side effects, a pharmaceutical composition formulated by combination of an angiotensin II-mediated compound or a salt thereof with at least one species of a compound having the activity of increasing insulin-sensitivity, a compound having the activity of improving postprandial hyperglycemia in diabetes mellitus, an indane derivative having the activity of inhibiting angiotensin converting enzyme, a pyridine derivative having the activity of inhibiting HMG-Co A reductase or salts thereof are advantageously employed.

DECLARATION AND POWER OF ATTORNEY

As a below named inventor, I hereby declare that:

My residence, post office address, and citizenship are as stated below next to my name.

I believe I am the original, first and sole inventor (if only one name is listed below) or an original, first and joint inventor (if plural names are listed below) of the subject matter which is claimed and for which a patent is sought on the invention entitled:

Pharmaceutical Composition,

the specification of which (check one)

☐ is attached hereto

☒ was filed on Apr. 3, 1997 as International Application No. PCT/JP97/01149 and was amended on _____ (if applicable).

I hereby state that I have reviewed and understand the contents of the above-identified specification, including the claims, as amended by any amendment referred to above.

I acknowledge the duty to disclose information which is known by me to be material to patentability as defined in Title 37, Code of Federal Regulations § 1.56.

I hereby claim foreign priority benefits under Title 35, United States Code, § 119 of any foreign application(s) for patent or inventor's certificate listed below and have also identified below any foreign application for patent or inventor's certificate having a filing date before that of the application on which priority is claimed:

PRIOR FOREIGN APPLICATION(S)

NUMBER	COUNTRY	DAY/MONTH/YEAR FILED	PRIORITY CLAIMED
83917-1996	Japan	05/04/1996	Yes

I hereby claim the benefit under Title 35, United States Code, § 120 of any United States application(s) listed below and insofar as the subject matter of each of the claims of this application is not disclosed in the prior United States application in the manner provided by the first paragraph of Title 35, United States Code, § 112, I acknowledge the duty to disclose information which is known by me to be material to patentability as defined in Title 37, Code of Federal Regulations § 1.56 which occurred between the filing date of the prior application and the national or PCT international filing date of this application:

APPLICATION SERIAL NO.	FILING DATE	STATUS: PATENTED, PENDING, ABANDONED

I hereby appoint as my attorneys, with full powers of substitution and revocation, to prosecute this application and transact all business in the Patent and Trademark Office connected therewith: Stephen A. Bent, Reg. No. 29,768; David A. Blumenthal, Reg. No. 26,257; John J. Feldhaus, Reg. No. 28,822; Donald D. Jeffery, Reg. No. 19,980; Eugene M. Lee, Reg. No. 32,039; Peter G. Mack, Reg. No. 26,001; Brian J. McNamara, Reg. No. 32,789; Sybil Meloy, Reg. No. 22,749; George E. Quillin, Reg. No. 32,792; Colin G. Sandercock, Reg. No. 31,298; Bernhard D. Saxe, Reg. No. 28,665; Richard L. Schwaab, Reg. No. 25,479; Arthur Schwartz, Reg. No. 22,115; Harold C. Wegner, Reg. No. 25,258.

Send all correspondence to FOLEY & LARDNER, Washington Harbour, 3000 K Street, N.W., Suite 500, P.O. Box 25696, Washington, D.C. 20007-8696. Address telephone communications to _____ at (202) 672-5300.

I hereby declare that all statements made herein of my own knowledge are true and that all statements made on information and belief are believed to be true; and further that these statements were made with the knowledge that willful false statements and the like so made are punishable by fine or imprisonment, or both, under Section 1001 of Title 18 of the United States Code and that such willful false statements may jeopardize the validity of the application or any patent issued thereon.

Full Name of First or Sole Inventor TAMURA Norikazu	Signature of First or Sole Inventor <i>Norikazu Tamura</i>	Date <i>May 8, 1997</i>
Residence Address 2-1-214, Ohgi 2-chome, Higashinada-ku, Kobe-shi HYOGO 658 JAPAN		Country of Citizenship Japan
Post Office Address Same as above		

Signatures should conform to names as typewritten. ☐ Additional inventors on attached Page 2.

Full Name of Second Inventor SOHDA Takashi	Signature of Second Inventor <i>Takashi Sohma</i>	Date May 8, 1997
Residence Address 27-20, Higashikanmaki 2-chome, Takatsuki-shi OSAKA 569 JAPAN	Country of Citizenship Japan	
Post Office Address Same as above		

Full Name of Third Inventor IKEDA Hitoshi	Signature of Third Inventor <i>Hitoshi Ikeda</i>	Date May 8, 1997
Residence Address 3-13-712, Nishiiwata 3-chome, Higashiosaka-shi OSAKA 578 JAPAN	Country of Citizenship Japan	
Post Office Address Same as above		

Full Name of Fourth Inventor	Signature of Fourth Inventor	Date
Residence Address	Country of Citizenship	
Post Office Address		

Full Name of Fifth Inventor	Signature of Fifth Inventor	Date
Residence Address	Country of Citizenship	
Post Office Address		

Full Name of Sixth Inventor	Signature of Sixth Inventor	Date
Residence Address	Country of Citizenship	
Post Office Address		

Full Name of Seventh Inventor	Signature of Seventh Inventor	Date
Residence Address	Country of Citizenship	
Post Office Address		